sanofi

MAINE RESIDENTS ONLY
APPLICATION

PATIENT Connection®

MAINE INSULIN SAFETY NET PROGRAM for SANOFI Insulins

The Act to Create the Insulin Safety Net Program ("the Act"), effective as of March 1, 2022, mandates that insulin manufacturers create an Insulin Safety Net Program ("the Program") to provide Maine (ME) residents who meet all eligibility criteria with their insulin prescription at no cost. Sanofi is administering the ME Insulin Safety Net Program through Sanofi Patient Connection®, a patient assistance program that helps patients get access to their medications.

What are the eligibility criteria mandated by the Act?

In order to be eligible for this Program, Maine requires you to meet the following requirements:

- You must be a resident of the State of Maine (ME) with a valid Maine identification card that indicates Maine residency in the form of a Maine identification card, driver's license, or permit. If the individual is under the age of 18, the individual's parent or legal guardian shall provide proof of residency.
- You must not be enrolled in MaineCare.
- You must not be enrolled in prescription drug coverage through an individual or group health plan that limits the total amount of cost-sharing that an enrollee is required to pay for a 30-day supply of insulin, including co-payments, deductibles, or coinsurance of $75 or less, regardless of the type or amount of insulin needed.
- You must have an annual household income of ≤400% of the current Federal Poverty Level.
- You must not be eligible to receive healthcare coverage through a federally funded program or receive prescription drug benefits through the United States Department of Veterans Affairs.
- If you are enrolled in Medicare Part D, in addition to the criteria above, you must also spend at least $1,000 on prescription drugs in the current calendar year.

How do I apply?

To apply for Patient Assistance Connection, all information must be complete and include the following:

Patient Information:

- Complete all relevant information on page 2, and sign and date the REQUIRED patient authorizations for HIPAA consent and income verification on page 2.

Healthcare Provider:

- Ask your healthcare provider (HCP) to complete page 3 and sign and date it.
- Ask your HCP to mail, fax, or submit through the Provider Portal your completed application (only pages 2 and 3 are needed).

Provide proof of ME residency (copy of ME ID card, driver's license, or permit). If you are under 18, your parent or legal guardian needs to provide residency proof. Missing information may delay processing of your application. Your completed application should be submitted by your HCP via mail or fax, including your completed application and residency proof to:

US Mail
Sanofi Patient Connection
PO Box 222138
Charlotte, NC 28222-2138

Fax
1.888.847.1797

What happens next?

Once we receive your application and proof of ME residency:

1. We will notify you within 5-7 business days if we require additional information to process your application.
2. Once we have all of the required information, we will review it to see if you meet all of ME's eligibility criteria to qualify for the Program within 3 business days.

If you are eligible:

1. You and your HCP will receive a letter within 10 business days notifying you of enrollment.
2. You will be enrolled for 12 months. If you are a Medicare Part D patient, you will be enrolled through the end of the calendar year. Your eligibility is renewable upon a redetermination of eligibility.
3. **Your medication will be sent directly to your home in approximately 5-7 business days from when you are approved.**

If you do not qualify for the Program, we will send you and your HCP a letter within 10 business days with the reason for denial.

DO NOT INCLUDE PATIENT MEDICAL RECORDS WITH THIS APPLICATION.

P: 1.888.847.4877 · F: 1.888.847.1797
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PATIENT TO FILL OUT

Section 1. Patient Information

Patient first name	MI	Last name
SSN		DOB
Address		City
State ME	Zip	Preferred language (if not English)
Phone number ( )		
Email		
Household size ○1 ○2 ○3 ○4 ○5 ○Other:		Annual household income

I permit Sanofi Patient Connection® to speak with the following person and/or organization about the information on this application and the status of my application request.

Patient representative/organization name | Relationship to patient | Phone

Patient Authorizations

Sanofi Patient Connection does **not** charge any fees for this service; application processing, medication, and shipping are all offered at no cost. Any fees charged to you by a third party completing this application on your behalf are not required by nor remitted to Sanofi.

I have read and agree to the HIPAA Consent included in **Section 6** on page 4.

PATIENT SIGN (REQUIRED)

(1 of 3) **Patient signature/Legal representative if patient is <18 years** Date

I have read and agree to the Income Verification included in **Section 7** on page 5.

PATIENT SIGN (REQUIRED)

(2 of 3) **Patient signature/Legal representative if patient is <18 years** Date

I have read and agree to the Patient Certifications regarding receiving communications from Sanofi Patient Connection included in **Section 8** on page 5.

PATIENT SIGN (OPTIONAL)

(3 of 3) **Patient signature/Legal representative if patient is <18 years** Date

Section 2. Insurance Information

Insurance? ○Yes ○No ☐If yes, is it Medicare Part D?

Primary insurance name		**Secondary** insurance name	
Insurance phone ( )		Insurance phone ( )	
Policy ID #	Group #	Policy ID #	Group #
Policyholder name (first/last)		Policyholder name (first/last)	
Relationship to patient	DOB	Relationship to patient	DOB

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PRESCRIBER TO FILL OUT

Please check the appropriate box (prescriber and patient signature required for all applications)

☐ **Patient Assistance**
No cost medication program. Check this box for no cost medication.

☐ **Benefits Verification (BV) and Patient Assistance**
Insurance coverage research and no cost medication program. Check this box if the patient has insurance coverage.

Section 3. Treatment and Prescribing Information *(See Section 5 for supported products)*

Patient name		DOB	
Concurrent medications			
Product #1	◯ Vials ◯ Pens ◯ N/A	Product #2	◯ Vials ◯ Pens ◯ N/A
ICD-10 Code		ICD-10 Code	
Frequency		Frequency	
Dosage (# of units per day)	Qty	Dosage (# of units per day)	Qty
If required by applicable state-specific law, please attach copies of prescriptions on official state prescription forms.			

Section 4. Prescriber Information

Prescriber name		State where licensed	
Site/facility name		Office contact email	
Type ◯ Clinic ◯ Physician office ◯ Outpatient hospital ◯ Inpatient hospital			Phone ( )
Facility address*			Fax ( )
City		State	Zip Code
Collaborating Physician Name			
License #	NPI #	Tax ID #	DEA #

**Sanofi product must be shipped to the signing prescriber's office or hospital address authorized by the prescriber and not to a third party.*

I certify that the information provided is current, complete, and accurate to the best of my knowledge. I certify that the Sanofi product is medically necessary for this patient and that I am authorized under state law to prescribe and dispense the requested medication. I certify that I have obtained from my patient all required written authorization for the release of my patient's personal identification, medical, and insurance information to Sanofi US and/or Sanofi Cares North America and their agents and representatives. I understand that any information provided is for the sole use of the Program to verify my patient's insurance coverage, to assess, if applicable, patient's eligibility for participation in the Patient Assistance Program and to otherwise administer the Sanofi Patient Connection Program and related services. I understand that I am under no obligation to prescribe any Sanofi product and that I have not received, nor will I receive, any benefit from Sanofi or their agents or representatives for prescribing a Sanofi product. The facility address noted above in Section 4 is my office or hospital address. My signature certifies that any prescription products received from this Program will be used for the above-named patient only and will not be resold nor offered for sale, trade, or barter and will not be returned for credit, nor will payment be sought from any payer, patient, or other source for product received from the Program.

HCP SIGN (REQUIRED)

Prescriber signature (REQUIRED – no stamps) **Printed name** **Date**

Section 5. Products Available With Patient Connection

- Admelog® (insulin lispro injection) 100 Units/mL
- Apidra® (insulin glulisine injection) 100 Units/mL
- Lantus® (insulin glargine injection) 100 Units/mL
- Soliqua® 100/33 (insulin glargine & lixisenatide) injection 100 Units/mL and 33 mcg/mL
- Toujeo® (insulin glargine injection) 300 Units/mL (1.5 mL or 3.0 mL pens)*

Full US prescribing information for all Sanofi Patient Connection supported products can be accessed at www.sanofipatientconnection.com/medications-available.
*Regular SoloStar® is packaged as 3 pens per pack, 450 units/pen; dials up to 80 units per single injection. Max SoloStar® is packaged as 2 pens per pack, 900 units/pen; dials up to 160 units per single injection; Max pen dials in 2-unit increments. Sanofi Patient Connection will provide assistance for any medically appropriate use as described in the prescribing information.

Additional Information

- Sanofi Patient Connection ships most medications in a 90-day supply.
- A representative from Sanofi may contact you for follow-up on any adverse event you may report regarding a Sanofi product.

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Section 6. Authorization to Use and Disclose Health Information (REQUIRED)

***Patient:* Please read the following carefully, then date and sign where indicated in Section 1 on page 2.**

HIPAA Consent: I authorize my healthcare providers and staff; my health insurer, health plan, or programs that provide me health benefits (together, "Health Insurers") to disclose to, Sanofi US, its affiliated companies (ie, Sanofi Pasteur U.S. and Genzyme, a Sanofi Company), Sanofi Cares North America, and authorized third-party agents involved in administration of this Program, (collectively "Program Sponsor"), health information about me, including information related to my medical condition, treatment, health insurance coverage, claims, prescriptions, and referral to and enrollment in this Program for purposes of determining my participation in, and administering, the Program, which may include contacting me as well as my doctor/healthcare provider, office/hospital staff, insurer (public/private) or others. I understand a representative from Sanofi may contact me for follow-up on any adverse event I may report regarding a Sanofi product. I authorize and consent to release of identifiable information about me including medical, financial and insurance records and information as required for participation in the Program. I understand that identifiable information about me will be kept confidential and will not be further used or disclosed except to administer the Program, or as required by law. I understand that information I authorize to be disclosed may be re-disclosed and is no longer protected by Federal privacy regulations. I agree that this authorization is voluntary and that I may refuse to sign this authorization. Refusal to sign will not affect my ability to obtain treatment but I will not be able to participate in this Program. Unless revoked, this authorization shall remain in effect throughout my participation in the Program, including subsequent reapplication as required. I may withdraw this authorization at any time by written notification to my doctor/healthcare provider; however, withdrawal of authorization will terminate my participation in this Program and will not affect information already disclosed under this Authorization.

I understand that it is my responsibility to follow up with my prescriber or the Program to make sure that my re-orders, as appropriate, are requested in a timely manner by my healthcare provider so I do not run out of medication. I understand that Sanofi US and Sanofi Cares North America reserve the right at any time and without notice to modify or change eligibility criteria or discontinue this Program. I understand that I may withdraw (take back) this Authorization at any time by calling 1.888.847.4877.

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Section 7. Income Verification

***Patient:* Please read the following carefully, then date and sign where indicated in Section 1 on page 2.**

Income Verification: I authorize Sanofi Patient Connection (SPC) 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, SPC 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 SPC 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 SPC 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. Continuation in the SPC Patient Assistance Program is conditioned upon timely verification of income. In addition, I agree to notify SPC if my insurance situation changes.

Section 8. Patient Certifications

***Patient:* Please read the following carefully, then date and sign where indicated in Section 1 on page 2.**

I authorize the Program to contact me by mail, telephone, or email, with information about the Program, disease state, and products, promotions, services, and research studies, and to ask my opinion about such information and topics, including market research and disease-related surveys. I further authorize the Program to de-identify my health information and use it in performing research, including linkage with other de-identified information the Program receives from other sources, education, business analytics, marketing studies, or for other commercial purposes. I understand that entities operating or administering parts of the Program may share identifiable health information with one another in order to de-identify it for these purposes and as needed to perform the Services or to send the communications listed above (the "Communications"). I understand and agree that the Program may use my health information for these purposes and may share my health information with my doctors, specialty pharmacies, and insurers. I understand that I may be contacted by the Program in the event that I report an adverse event associated with a Sanofi product.

I understand that I do not have to opt in to receive the Communications, and that I can still receive patient assistance through the Program, as prescribed by my physician. I may opt out of receiving the Communications offered by the Program, at any time by notifying a Program representative by telephone at 1-800-633-1610 or by mailing a letter to Sanofi US Customer Services, P.O. Box 5925 Mailstop 55A-220A5, Bridgewater, NJ 08807-5925. I also understand that the Services may be revised, changed, or terminated at any time.

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