



**Division of Genomic Diagnostics
PROBAND EXOME/GENOME REQUISITION AND
CONSENT FOR CHOP PATIENTS**

Page 1 of 7

LAST NAME

FIRST NAME

MR#

DOB

 PLACE PATIENT LABEL HERE OR COMPLETE ABOVE

DO NOT HANDWRITE PATIENT INFORMATION HERE

SHIP TO: Children's Hospital of Philadelphia, Genomic Diagnostics Laboratory, 3615 Civic Center Blvd., Abramson Research Center, 714J, Philadelphia, PA 19104-4302 - Phone: (267) 426-1447

PROBAND INFORMATION		TEST INFORMATION
Patient Name (Last, First, Middle): _____ Ordering Facility MRN: _____ Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Unknown <input type="checkbox"/> Other: _____ Date of Birth (Month/Day/Year): _____ Street Address: _____ City/State: _____ Zip/Country: _____ Phone Number: (____) _____ Race/Ethnicity: <input type="checkbox"/> Amish <input type="checkbox"/> Asian <input type="checkbox"/> Black/African-American <input type="checkbox"/> Caucasian <input type="checkbox"/> East Indian <input type="checkbox"/> French Canadian <input type="checkbox"/> Hispanic <input type="checkbox"/> Jewish-Ashkenazi <input type="checkbox"/> Jewish-Sephardic <input type="checkbox"/> Mediterranean <input type="checkbox"/> Native American <input type="checkbox"/> Other: _____ Sample Type: <input type="checkbox"/> Peripheral Blood in EDTA tube > 3 mL <input type="checkbox"/> Saliva (Contact lab for kits) <input type="checkbox"/> Cord Blood in EDTA tube (Maternal Sample Required) <input type="checkbox"/> *DNA >15 µg Specify Tissue of Origin: _____ <input type="checkbox"/> Other: _____ <i>If interested in sending a specimen other than blood, saliva, or DNA, please call the lab to discuss.</i> <i>*Nucleic acids (DNA or RNA) must be extracted from a CAP or CLIA certified laboratory.</i> Collection Date (Month/Day/Year): _____ Collection Time: _____ AM/PM		Test Requested: <input type="checkbox"/> CHOP Medical Exome, Proband & Family <input type="checkbox"/> CHOP Medical Exome, Proband Only <input type="checkbox"/> CHOP Medical Exome Reanalysis, Proband & Family* <input type="checkbox"/> CHOP Medical Exome Reanalysis, Proband Only <input type="checkbox"/> CHOP Medical Exome from Panel <input type="checkbox"/> CHOP Medical Exome + MitoGenome Analysis <input type="checkbox"/> CHOP Rapid Medical Genome, Proband & Family <input type="checkbox"/> CHOP Rapid Medical Genome, Proband <input type="checkbox"/> CHOP Rapid Medical Genome from Panel <input type="checkbox"/> CHOP Medical Genome, Proband & Family <input type="checkbox"/> CHOP Medical Genome, Proband Only <input type="checkbox"/> CHOP Medical Genome Reanalysis, Proband & Family* <input type="checkbox"/> CHOP Medical Genome Reanalysis, Proband Only <input type="checkbox"/> CHOP Medical Genome from Panel <small>NOTE: For any of the tests above, additional family member specimens may be submitted for Sanger sequencing to assist with interpretation. If family member specimens are to be submitted, please indicate this on the next page.</small> <small>*Please note: New family members (who did not undergo exome/genome analysis with the original order) can only be submitted for Sanger sequencing.</small> Additional Genetic Testing: <input type="checkbox"/> Patient has additional genetic testing ordered on the General Test Requisition Checklist of Items to Include: <input type="checkbox"/> Proband specimen <input type="checkbox"/> Specimen of proband's mother (if applicable) <input type="checkbox"/> Specimen of proband's father (if applicable) <input type="checkbox"/> Specimens of other family members to be analyzed (please contact lab to discuss) <input type="checkbox"/> Test requisition form (all billing and clinical information must be completed) <input type="checkbox"/> Signed informed consent form <input type="checkbox"/> Family history and pedigree <input type="checkbox"/> Proband medical records and photographs
ORDERING PROVIDER	ORDERING LABORATORY	OTHER ORDERING PROVIDER / GENETIC COUNSELOR
Name (Last, First, Degree) (____) _____ Phone (____) _____ Fax Institution Street Address City State Zip Country Email	Name (Last, First, Degree) (____) _____ Phone (____) _____ Fax Institution Street Address City State Zip Country Email	Name (Last, First, Degree) (____) _____ Phone (____) _____ Fax Email

For Lab Use Only
 Type of billing: Institutional CHOP Self-pay Institution called
 Comments:

Received by: _____ Received Date: _____ Received Time: _____ Sample: _____

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**Division of Genomic Diagnostics
PROBAND EXOME/GENOME REQUISITION AND
CONSENT FOR CHOP PATIENTS**

Page 2 of 7

SPECIMEN INFORMATION FOR PARENTS AND OTHER FAMILY MEMBERS (IF APPLICABLE)

Proband's Mother's Full Name: _____

Proband's Mother's Date of Birth: _____ Proband's Mother's MRN: _____
Month/Day/Year

Considering the proband's phenotype, the mother is: Affected Unaffected Unknown

Mother's specimen: Included with this test requisition form To be sent later Not available

Family Member Test requested to aid in proband exome/genome analysis data interpretation: Exome/Genome analysis
 Sanger confirmation only

Proband's Father's Full Name: _____

Proband's Father's Date of Birth: _____ Proband's Father's MRN: _____
Month/Day/Year

Considering the proband's phenotype, the father is: Affected Unaffected Unknown

Father's specimen: Included with this test requisition form To be sent later Not available

Family Member Test requested to aid in proband exome/genome analysis data interpretation: Exome/Genome analysis
 Sanger confirmation only

Family Member's Full Name: _____ Relationship to Proband: _____

Family Member's Date of Birth: _____ Family Member's MRN: _____
Month/Day/Year

Considering the proband's phenotype, the family member is: Affected Unaffected Unknown

Family Member's specimen: Included with this test requisition form To be sent later Not available

Family Member Test requested to aid in proband exome/genome analysis data interpretation: Exome/Genome analysis
 Sanger confirmation only

Family Member's Full Name: _____ Relationship to Proband: _____

Family Member's Date of Birth: _____ Family Member's MRN: _____
Month/Day/Year

Considering the proband's phenotype, the family member is: Affected Unaffected Unknown

Family Member's specimen: Included with this test requisition form To be sent later Not available

Family Member Test requested to aid in proband exome/genome analysis data interpretation: Exome/Genome analysis
 Sanger confirmation only

AII SPECIMENS MUST BE CLEARLY LABELED WITH FULL NAME AND DATE OF BIRTH

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CONSENT FOR CHOP PATIENTS

Page 3 of 7

PROBAND EXOME/GENOME ANALYSIS CONSENT FORM

This form was designed to guide the consent process for exome analysis or genome analysis and help you to have a discussion with your healthcare provider. Your healthcare provider will help determine which test is most appropriate for your circumstances. You can decide whether or not you want to have this test. We encourage you to ask questions, including questions about other testing options.

What are Exome Analysis and Genome Analysis?

- **Exome analysis and genome analysis** are genetic tests that examine most of a person's genes all at once. **Genes** tell our body how to grow and develop.
- These tests can find differences in genes that may explain a patient's health and/or developmental concerns. These genetic differences may also be called **variants** or **alterations**. The primary patient having testing may also be called the **proband**. Genome analysis may identify and report variants that exome analysis does not.
- For more information about exome analysis, genome analysis and genetic testing, visit the website: <https://imgc.chop.edu/types-of-genetic-testing/>.

How are these tests performed at CHOP?

- These tests require a biological sample, such as blood, from the patient and from one or more biological (or blood-related) family members (if available). The laboratory will obtain and look at the DNA (genetic material) from the sample(s).
- The laboratory will compare the patient's DNA sequence (or pattern) to the sequence usually found in healthy people, as well as to that of other family members (if applicable). There may be thousands of differences between the DNA sequences. Most of these differences are normal variation and do not cause health problems.
- The laboratory will use information about the patient's health and the health of family members to identify whether any of the many DNA differences may be the cause of the patient's reason for testing.
- The laboratory will inform the referring healthcare provider(s) about the findings that are most likely to be related to the patient's **reason for testing** or **clinical indication**.

Clinical Information and Testing of Family Members

- The laboratory needs to have correct information about the health of the patient and family members in order to interpret the results from these tests properly.
- The laboratory will use the reason for testing provided by the healthcare provider(s), as well as additional medical records/history, in order to interpret the results. The laboratory does not review the entirety of the patient's medical record/history.
- The laboratory is more likely to identify the genetic cause of a patient's condition when relevant family members are tested at the same time.

Secondary Findings

- The American College of Medical Genetics and Genomics (ACMG) recommends that laboratories performing exome or genome analysis *purposefully* seek and report findings that cause a specific group of rare conditions, even if those conditions are not the reason for the patient's testing. These results are called **secondary findings** because they are not related to the patient's current reason for testing.
- The conditions associated with secondary findings may lead to serious health problems, such as an increased risk of cancer or heart rhythm issues, or an increased risk for complications from certain kinds of anesthesia. These conditions may be inherited. Some may cause symptoms in infancy or childhood, while others do not usually cause symptoms until adulthood.
- These conditions can in some instances be improved or avoided when monitored or treated. Therefore, they are considered **medically actionable**.
- Depending on the age and sex of the patient, some secondary findings may have a specific change in medical management that could be pursued within the patient's near (immediate) future and lead to a clear improvement in health outcomes. These findings are considered **immediately medically actionable secondary findings** in the patient.

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Division of Genomic Diagnostics
PROBAND EXOME/GENOME REQUISITION AND
CONSENT FOR CHOP PATIENTS

Page 4 of 7

- A full Secondary Findings List detailing the genes and conditions currently included in the secondary findings analysis for this test is available on the laboratory website at <http://www.chop.edu/centers-programs/division-genomic-diagnostics/requisition-forms> or by contacting the laboratory at 267-426-1447.
- **Options for Secondary Findings:**
 - **Testing for secondary findings is optional, and you will make a selection at the end of this form.**
 - If you select **Option 1** below, the laboratory will purposefully look for and report ALL secondary findings that are likely to cause disease. It is possible that these tests will not detect all secondary findings, even if Option 1 is selected. In addition, these tests will not look for all causes of the conditions on the Secondary Findings List.
 - If you select **Option 2** below, the laboratory will not purposefully look for or report ANY findings on the Secondary Findings List, even if they are identified *by chance*, unless the finding is possibly related to the patient's reason for testing.
 - If you select **Option 3** below, the laboratory will not purposefully look for findings on the Secondary Findings List, but will report findings from the list that are identified *by chance* and are *immediately medically actionable* in the patient or that are possibly related to the patient's reason for testing.
 - If you select **Option 2 or 3**, unreported secondary findings will only be available in the future if a formal exome reanalysis or genome reanalysis is requested.
- **Secondary Findings in Family Members:**
 - If a secondary finding is identified in the patient, the laboratory can also look for the finding in family members who have exome analysis or genome analysis at the same time as the patient.
 - Family members select whether or not they want to receive this information on the Family Member Test Requisition and Consent form.

Other Findings Unrelated to the Reason for Testing

- In addition to secondary findings, these tests may in some rare instances detect other results unrelated to the patient's reason for testing. Unlike the secondary findings described above, these other unrelated findings are NOT purposefully sought but are found by chance while performing these tests. These findings are sometimes called **incidental findings** because they are found incidentally.
- The lab will report **medically actionable incidental findings** that are likely to cause serious health problems and have a well-established change in management or treatment. These include childhood onset conditions or conditions with a recommended change in medical care that could take place within about ten or twenty years.
- We generally only report incidental findings that are medically actionable. There are rare exceptions when we may report findings that are *not* medically actionable. Examples include but are not limited to: 1) if they explain some of the patient's symptoms that are not included in the clinical indication for testing, 2) if they are associated with another finding (such as a large deletion or missing region of genetic material) that is believed to be associated with the patient's clinical indication for testing, or 3) if they are associated with an untreatable, serious, childhood onset disorder that is highly likely to cause symptoms (as reporting these findings could help to avoid additional testing and future delays in diagnosis).

What will the Laboratory Find and Report?

- The test result might be **diagnostic, inconclusive, or negative**.
 - A **diagnostic** result means that the test identified a genetic variant that is likely to explain the patient's reason for testing.
 - An **inconclusive** result means that the test identified variant(s) in a gene that may explain the patient's reason for testing, but there is not enough information available to be certain. (The laboratory will inform the patient's healthcare provider if a clinically significant change in interpretation is identified in the future.)
 - A **negative** result means that the test did not identify any genetic variants that could explain the patient's reason for testing. This result does not rule out the possibility that the patient has a genetic disorder that was unable to be detected by the exome or genome test at this time.

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Division of Genomic Diagnostics
PROBAND EXOME/GENOME REQUISITION AND
CONSENT FOR CHOP PATIENTS

Page 5 of 7

- The report will include:
 - DNA variants that were detected that may be the cause of the patient's reason for testing and which family members have this variant (when possible).
 - DNA variants that were detected and are *not* related to the patient's reason for testing and which family members have this variant (when possible):
 - 1) Secondary findings, only if Option 1 below is selected.
 - 2) Incidental findings identified by chance that are medically actionable or meet the exception criteria above.
- All findings will be included in the patient's report. Separate reports will *not* be issued automatically for family members.

What are the Limitations of Exome Analysis and Genome Analysis?

- These tests do not look at all of a person's genes. Many genes will not be examined completely or at all.
- These tests do not find all genetic variants or risk factors for common diseases.
- These tests do not identify all types of DNA variants.
- The results usually do not predict how serious a health condition will be or at what age it will occur.
- The results may or may not improve the patient's treatment or prognosis.
- These tests will not always identify the cause of a patient's reason for testing. We expect these tests to identify the cause of a patient's reason for testing in about 1 out of every 4 (25%) patients who are tested.
- Exome analysis and genome analysis and our interpretation of the results are based on what is known about genetic diseases at the time the test is performed. As time passes, more information about genetic diseases and genetic testing may become known. Exome and genome tests performed in the future with new knowledge may provide different results. Although the laboratory may perform limited reanalysis of exome and genome data for internal purposes (such as quality improvement), laboratory-initiated reanalysis of the raw exome and genome data is not performed routinely. **A healthcare provider may request reanalysis of exome and genome data as a part of future care for an additional fee.** A new consent form may not be requested if your provider orders reanalysis.

What are the Potential Risks of Exome Analysis and Genome Analysis?

- These tests could suggest an incorrect genetic cause for the patient's symptoms or not identify the true cause of a patient's reason for testing, due to limitations of current scientific knowledge and technology.
- We may learn more about these results in the future. The results you receive from the test may later be interpreted differently as a result of evolving research and knowledge in the field. It is appropriate to follow up with your clinician from time to time regarding possible new information about the test results.
- These tests may identify results that have an uncertain meaning (variants of uncertain significance).
- These tests may reveal unexpected findings unrelated to the patient's reason for testing.
- Although it is not the intended purpose, these tests may identify a difference in biological family relationships that may or may not be expected by the family. For example, the results may show that the patient's biological mother or father is someone different than originally believed. It may be possible to tell a difference in biological family relationships from the results report. In some cases, such as when results suggest the possibility of incest, the hospital may be required to report this information to appropriate authorities.
- These tests may find results that impact the health or future reproductive decisions of the patient or family members.
- These tests may find results that could affect the patient's or family member's ability to buy life insurance, disability insurance, or long-term care insurance in the future.
- Because exome analysis and genome analysis are complex, the cost of these tests may be higher than other genetic testing. Exome analysis and/or genome analysis may or may not be covered by insurance.
- If you are concerned about cost, please discuss your questions with your healthcare provider, insurance company, or the CHOP Family Health Coverage Program (1-267-426-0359) before proceeding with testing.

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CONSENT FOR CHOP PATIENTS

Page 6 of 7

Results Reporting and Confidentiality

- We will report the results to the healthcare provider(s) or institution who ordered the test.
- We will release results to individuals not associated with CHOP if we have written permission from the patient, the patient's legal representative, or as required by law.
- The results report will become part of the patient's medical record and therefore has the potential for future release.
- The patient's health insurance provider or other parties may have legal access to the test results.
- In rare instances, CHOP may ask external (non-CHOP) clinical laboratories to perform all or part of the exome or genome test. The external laboratory may keep copies of the patient's genetic information, including raw genetic data and/or results reports.
- For more information about how CHOP can use or share your information see the Notice of CHOP Privacy Practices (https://at.chop.edu/general-counsel/compliance-privacy/privacy/Shared%20Documents/hipaa_npp.pdf).
- The samples and raw genetic data relating to the patient without identifiers, such as name and date of birth, may be maintained in CHOP-based databases and may be shared with external genetic databases that are not located at, owned by, or operated by CHOP. These databases were created to improve our understanding and interpretation of genetic results and ensure that clinical genetic laboratories are interpreting results in the same way. The information contained in these databases may also be used for research purposes.

Future of the Information and Samples

- **Storage and Sharing of the Information and Samples**
 - We will keep raw genetic data for at least two years, as per professional and regulatory guidelines. We may destroy it after that time.
 - CHOP is not a DNA storage or banking facility. There is no guarantee that patient samples will be available or usable for additional or future testing.
 - We will keep the patient's raw genetic data on a secure server/computer that is in a separate section of the medical record and only accessible by authorized personnel. Future access to this data may or may not be available. If you want access to the raw genetic data, please contact the laboratory. There may be an associated cost.
 - The laboratory will only release the raw genetic data with identifiers, such as name or date of birth, with the written permission of the patient, patient's legal representative, or as required by law.
 - External clinical laboratories who perform all or part of the exome or genome test may keep copies of the patient's genetic information.
 - The samples and raw genetic data relating to the patient without identifiers, such as name and date of birth, may be maintained in CHOP-based databases and may be shared with external genetic databases that are not located at, owned by, or operated by CHOP. These databases were created to improve our understanding and interpretation of genetic results and ensure that clinical genetic laboratories are interpreting results in the same way. The information contained in these databases may also be used for research purposes.
- **Reanalysis**
 - A healthcare provider may request reanalysis of exome and genome data as a part of future care for an additional fee.
 - In addition, although not performed routinely, the laboratory may independently initiate limited reanalysis of a patient's (and any submitted family members') exome or genome data for internal purposes, such as quality improvement. We may share new relevant information from this laboratory-initiated reanalysis with your ordering provider or other appropriate healthcare providers. Please discuss with your healthcare provider about receiving any updated information.

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**Division of Genomic Diagnostics
PROBAND EXOME/GENOME REQUISITION AND
CONSENT FOR CHOP PATIENTS**

Page 7 of 7

Choice for Secondary Findings

Please Note: the laboratory cannot proceed with the test until you select an option below

Please initial one of the options below regarding your choice for secondary findings in the patient. (The adult patient or a parent/guardian of a minor child must initial next to the selected option.)

Secondary Findings Options				
Initial in the Box Next to Your Choice	Option	Lab Looks for Secondary Findings	Lab Reports Secondary Findings	Details
	Option 1	Yes	Yes	I want the lab to look for and report ALL detected disease causing variants on the Secondary Findings List. NOTE: It is possible that this test will not detect all secondary findings. In addition, this test will not look for all causes of the conditions on the Secondary Findings List.
	Option 2	No	No	I DO NOT want the lab to look for or report <i>any</i> secondary findings. I understand that some of these findings may be <i>immediately medically actionable</i> in the patient. I am aware that I will not have access to these results later. NOTE: If you choose this option, findings that are possibly related to the patient's reason for testing and also involved a gene/condition on the Secondary Findings List will still be reported.
	Option 3	No	Yes, if found by chance and immediately medically actionable	I DO NOT want the lab to purposefully look for secondary findings. I want the lab to report secondary findings identified by <i>chance</i> if they are <i>immediately medically actionable</i> in the patient.

Genetic Counseling and Clinical Interpretation

- We recommend that you seek genetic counseling before you decide whether or not you want to have this test and when you receive the test results.
- If you would like to see a genetic counselor, please ask your healthcare provider to refer you to one in your area. You can find a genetic counselor near you by visiting www.nsgc.org.
- The healthcare provider who ordered this test will make the final interpretation about what the results mean for the patient and provide follow-up recommendations.

By signing this document, you are agreeing that the test, its limitations, and use and retention of related data and samples have been explained to you and you consent and agree to the test and these uses.

Patient/Parent/Guardian Statement: I acknowledge that I have discussed the benefits, risks, and limitations of exome/genome analysis with my healthcare provider(s). I consent to testing.

Patient Signature

Printed Name

Date

Time

Parent/Legal Guardian if Different from Patient Signature

Printed Name

Date

Time

Healthcare Provider's Statement: I have explained the benefits, risks, limitations, and data use and retention of exome analysis and/or genome analysis to this individual and addressed his/her questions about the test(s). I understand that it is my responsibility to interpret the clinical relevance of the results for this individual and to provide appropriate follow-up recommendations.

Healthcare Provider Signature

Printed Name

and Contact Number

Date

Time

Interpreter Signature/Witness Signature (Circle Relevant Role)

Printed Name

Date

Time

Phone: (267) 426-1447; Fax: (215) 590-3514; DGDGeneticCounselor@chop.edu; www.chop.edu/centers-programs/division-genomic-diagnostics

*Testing will not be initiated until all necessary samples and clinical information are submitted to the laboratory.

OUR COMMITMENT TO DIVERSE POPULATIONS

Children's Hospital of Philadelphia complies with applicable federal civil rights laws and does not discriminate on the basis of race, color, national origin, age, disability or sex. Children's Hospital of Philadelphia does not exclude people or treat them differently because of race, color, national origin, age, disability or sex.

CHILDREN'S HOSPITAL OF PHILADELPHIA:

- Provides reasonable modifications and free aids and services to help people with disabilities to communicate effectively with us, such as:
 - Qualified sign language interpreters
 - Written information in other formats (large print, audio, accessible electronic formats, other formats)
- Provides free language services to people whose primary language is not English, such as:
 - Qualified interpreters
 - Information written in other languages

If you need these services, call 800-879-2467.

If you believe that Children's Hospital of Philadelphia has failed to provide these services or discriminated in another way on the basis of race, color, national origin, age, disability, sex, or any other protected classification, you can contact the Section 1557 Coordinator (1557coordinator@chop.edu) or file a grievance with:

The Family Relations Office
3401 Civic Center Blvd., Philadelphia, PA 19104
Phone: 267-426-6983

- email: familyrelations@chop.edu.

You can file a grievance in person or by mail, or email.

If you need help filing a grievance, Family Relations is available to help you. You can also file a civil rights complaint with the U.S. Department of Health and Human Services, Office for Civil Rights, electronically through the Office for Civil Rights Complaint Portal, available at ocrportal.hhs.gov/ocr/portal, or by mail or phone at:

U.S. Department of Health and Human Services
200 Independence Ave. • SW Room 509F, HHH Building • Washington, DC 20201
Phone: 800-368-1019; 800-537-7697 (TDD)

Complaint forms are available at hhs.gov/ocr/office/file/index.html.

This notice is available at <https://www.chop.edu/nondiscrimination-statement>

Notice of Availability of Language Assistance Services and Auxiliary Aids and Services (§ 92.11)

English

ATTENTION: If you speak, free language assistance services are available to you. Appropriate auxiliary aids and services to provide information in accessible formats are also available free of charge. Call 1-800-879-2467, then press 1 or speak to your provider.

Español - Spanish

ATENCIÓN: Si habla español, tiene a su disposición servicios gratuitos de asistencia lingüística. También están disponibles de forma gratuita ayuda y servicios auxiliares apropiados para proporcionar información en formatos accesibles. Llame al 1-800-879-2467, luego presione 1 o hable con su proveedor.

中文- Chinese

注意：如果您说中文，我们将免费为您提供语言协助服务。我们还免费提供适当的辅助工具和服务，以无障碍格式提供信息。致电 1-800-879-2467 然后按 1 或咨询您的服务提供商。

Arabic
العربية
تنبيه: إذا كنت تتحدث اللغة العربية، فستتوفر لك خدمات المساعدة اللغوية المجانية. كما تتوفر وسائل مساعدة وخدمات مناسبة لتوفير المعلومات بتنسيقات يمكن الوصول إليها مجانًا. اتصل على الرقم 18008792467، ثم اضغط على الرقم ١.

РУССКИЙ - Russian

ВНИМАНИЕ: Если вы говорите на русский, вам доступны бесплатные услуги языковой поддержки. Соответствующие вспомогательные средства и услуги по предоставлению информации в доступных форматах также предоставляются бесплатно. Позвоните по телефону 1-800-879-2467 затем нажмите 1 или обратитесь к своему поставщику услуг.

Português - Portuguese

ATENÇÃO: Se você fala Português, serviços gratuitos de assistência linguística estão disponíveis para você. Auxílios e serviços auxiliares apropriados para fornecer informações em formatos acessíveis também estão disponíveis gratuitamente. Ligue para 1-800-879-2467 em seguida, pressione 1 ou fale com seu provedor.

Việt - Vietnamese

LƯU Ý: Nếu bạn nói tiếng Việt, chúng tôi cung cấp miễn phí các dịch vụ hỗ trợ ngôn ngữ. Các hỗ trợ dịch vụ phù hợp để cung cấp thông tin theo các định dạng dễ tiếp cận cũng được cung cấp miễn phí. Vui lòng gọi theo số 1-800-879-2467 sau đó nhấn 1 hoặc trao đổi với người cung cấp dịch vụ của bạn.

Français - French

ATTENTION : Si vous parlez Français, des services d'assistance linguistique gratuits sont à votre disposition. Des aides et services auxiliares appropriés pour fournir des informations dans des formats accessibles sont également disponibles gratuitement. Appelez le 1-800-879-2467 puis appuyez sur 1 ou parlez à votre fournisseur.

Kreyòl Ayisyen - Haitian-Creole

ATANSYON: Si w pale Kreyòl Ayisyen, gen sèvis èd aladispozisyon w gratis pou lang ou pale a. Èd ak sèvis siplemantè apwopriye pou bay enfòmasyon nan fòm aksesib yo disponib gratis tou. Rele nan 1-800-879-2467 apre sa peze 1 oswa pale avèk founisè w la.

Italiano - Italian

ATTENZIONE: se parli Italiano, sono disponibili servizi di assistenza linguistica gratuiti. Sono inoltre disponibili gratuitamente ausili e servizi ausiliari adeguati per fornire informazioni in formati accessibili. Chiama il 1-800-879-2467 poi premi 1 o parla con il tuo fornitore.

한국어 - Korean

주의: 한국어를 사용하시는 경우 무료 언어 지원 서비스를 이용하실 수 있습니다. 이용 가능한 형식으로 정보를 제공하는 적절한 보조 기구 및 서비스도 무료로 제공됩니다. 1-800-879-2467 그런 다음 1 을 누르세요 번으로 전화하거나 서비스 제공업체에 문의하십시오.

नेपाली - Nepali

नोट: यदि तपाईं नेपाली बोल्नुहुन्छ भने तपाईंलाई निःशुल्क भाषा समर्थन सेवाहरू उपलब्ध छन्। पहुँचयोग्य ढाँचाहरूमा जानकारी प्रदान गर्न उपयुक्त समर्थन र सेवाहरू पनि निःशुल्क उपलब्ध छन्। 1-800-879-2467 मा कल गर्नुहोस् र त्यसपछि 1 थिच्नुहोस् वा आफ्नो प्रदायकसँग कुरा गर्नुहोस्।

हिंदी - Hindi

ध्यान दें: यदि आप हिंदी बोलते हैं, तो आपके लिए निःशुल्क भाषा सहायता सेवाएं उपलब्ध होती हैं। सुलभ प्रारूपों में जानकारी प्रदान करने के लिए उपयुक्त सहायक साधन और सेवाएँ भी निःशुल्क उपलब्ध हैं। 1-800-879-2467 फिर 1 दबाएँ। पर कॉल करें या अपने प्रदाता से बात करें।

Deutsch - German

ACHTUNG: Wenn Sie Deutsch sprechen, stehen Ihnen kostenlose Sprachassistentendienste zur Verfügung. Entsprechende Hilfsmittel und Dienste zur Bereitstellung von Informationen in barrierefreien Formaten stehen ebenfalls kostenlos zur Verfügung. Rufen Sie 1-800-879-2467 an, dann drücken Sie die 1 oder sprechen Sie mit Ihrem Gesundheitsdienstleiter.

ગુજરાતી - Gujarati

ધ્યાન આપો: જો તમે ગુજરાતી બોલતા હો તો મફત ભાષાકીય સહાયતા સેવાઓ તમારા માટે ઉપલબ્ધ છે. યોગ્ય ઓક્રિઝલરી સહાય અને એક્સેસિબલ ફોર્મેટમાં માહિતી પૂરી પાડવા માટેની સેવાઓ પણ વિના મૂલ્યે ઉપલબ્ધ છે. 1-800-879-2467 પછી 1 દબાવી પર કોલ કરો અથવા તમારા પ્રદાતા સાથે વાત કરો.

Polski - Polish

UWAGA: Osoby mówiące po polsku mogą skorzystać z bezpłatnej pomocy językowej. Dodatkowe pomoce i usługi zapewniające informacje w dostępnych formatach są również dostępne bezpłatnie. Zadzwoń pod numer 1-800-879-2467 następnie naciśnij 1 lub porozmawiaj ze swoim dostawcą.

українська мова - Ukrainian

УВАГА: Якщо ви розмовляєте українською мовою, вам доступні безкоштовні мовні послуги. Відповідні допоміжні засоби та послуги для надання інформації у доступних форматах також доступні безкоштовно. Зателефонуйте за номером 1-800-879-2467 потім натисніть 1 або зверніться до свого постачальника.

Tagalog – Tagalog

PAALALA: Kung nagsasalita ka ng Tagalog, magagamit mo ang mga libreng serbisyong tulong sa wika. Magagamit din nang libre ang mga naaangkop na auxiliary na tulong at serbisyong upang magbigay ng impormasyon sa mga naa-access na format. Tumawag sa 1-800-879-2467 pagkatapos ay pindutin ang 1 o makipag-usap sa iyong provider.

తెలుగు - Telugu

సావధానం: మీరు తెలుగు మాట్లాడితే, మీకు ఉచిత భాషా సహాయ సేవలు అందుబాటులో ఉంటాయి. యాక్సెస్ చేయగల ఫార్మాట్లలో సమాచారాన్ని అందించడానికి తగిన సహాయక సహాయాలు మరియు సేవలు కూడా ఉచితంగా అందుబాటులో ఉంటాయి. 1-800-879-2467 తర్వాత 1 నొక్కండి కి కాలి చేయండి లేదా మీ ప్రావైడర్‌తో మాట్లాడండి.

Deitsch - Pennsylvania Dutch

Wann du Deitsch (Pennsylvania German/Dutch) schwetztscht, kannscht du mitaus Koschte ebber gricke, ass dihr helft mit die englisch Schprooch. G'geignete Hilfsmiddel un Dienschtför Information in zugängliche Formate sin aa fer nix do. Ruf 1-800-879-2467 aa, dann drick 1 oder schwetz mit dei Dienschtleistender.