MEDICAL POLICY – 2.01.31
Intra-Articular Hyaluronan Injections for Osteoarthritis

BCBSA Ref. Policy: 2.01.31
Effective Date: July 1, 2019
Last Revised: Oct. 1, 2019
Replaces: N/A
RELATED MEDICAL POLICIES:
7.01.549 Knee Arthroscopy in Adults

Select a hyperlink below to be directed to that section.

POLICY CRITERIA | CODING | RELATED INFORMATION
EVIDENCE REVIEW | REFERENCES | HISTORY

∞ Clicking this icon returns you to the hyperlinks menu above.

Introduction

Hyaluronan, or sodium hyaluronate, is a natural substance found in some joints of the human body. This substance acts as a lubricant to help the joints work better. There are also other substances similar to hyaluronan that have been used for the past several years to try to treat knee pain due to osteoarthritis of the knee. These substances have been manufactured to be injected into the knee one or more times. Early studies suggested the injections helped decrease pain. However, during the past 5 years, re-analysis of the best early medical studies that included thousands of patients have shown that these injections do not offer significant relief for most people. Some professional societies recommend that these injections not be used. Based on the changing scientific evidence, the plan considers hyaluronan injections in the knee not medically necessary. Use of these injections in all other joints is considered investigational (unproven) because there are so few studies published about other joints. The plan does not pay for services that are not medically necessary or investigational.

Note: The Introduction section is for your general knowledge and is not to be taken as policy coverage criteria. The rest of the policy uses specific words and concepts familiar to medical professionals. It is intended for providers. A provider can be a person, such as a doctor, nurse, psychologist, or dentist. A provider also can be a place where medical care is given, like a hospital, clinic, or lab. This policy informs them about when a service may be covered.
### Procedure

<table>
<thead>
<tr>
<th>Medical Necessity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intra-articular hyaluronan injections of the knee are considered not medically necessary.</td>
</tr>
</tbody>
</table>

### Procedure

<table>
<thead>
<tr>
<th>Investigational</th>
</tr>
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<tbody>
<tr>
<td>Intra-articular hyaluronan injections are considered investigational for all other joints.</td>
</tr>
</tbody>
</table>

### Coding

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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</thead>
<tbody>
<tr>
<td><strong>HCPCS</strong></td>
<td></td>
</tr>
<tr>
<td>J7318</td>
<td>Hyaluronan or derivative, durolane, for intra-articular injection, 1 mg (new code effective 1/1/19)</td>
</tr>
<tr>
<td>J7320</td>
<td>Hyaluronan or derivative, Genvisc 850, for intra-articular injection, 1 mg</td>
</tr>
<tr>
<td>J7321</td>
<td>Hyaluronan or derivative, Hyalgan or Supartz, for intra-articular injection, per dose</td>
</tr>
<tr>
<td>J7322</td>
<td>Hyaluronan or derivative, Hymovis, for intra-articular injection, 1 mg</td>
</tr>
<tr>
<td>J7323</td>
<td>Hyaluronan or derivative, Euflexxa, for intra-articular injection, per dose</td>
</tr>
<tr>
<td>J7324</td>
<td>Hyaluronan or derivative, Orthovisc, for intra-articular injection, per dose</td>
</tr>
<tr>
<td>J7325</td>
<td>Hyaluronan or derivative, Synvisc or Synvisc-One, for intra-articular injection, 1 mg</td>
</tr>
<tr>
<td>J7326</td>
<td>Hyaluronan or derivative, Gel-One, for intra-articular injection, per dose</td>
</tr>
<tr>
<td>J7327</td>
<td>Hyaluronan or derivative, Monovisc, for intra-articular injection, per dose</td>
</tr>
<tr>
<td>J7328</td>
<td>Hyaluronan or derivative, for intra-articular injection, 0.1 mg</td>
</tr>
<tr>
<td>J7329</td>
<td>Hyaluronan or derivative, trivisc, for intra-articular injection, 1 mg (new code effective 1/1/19)</td>
</tr>
<tr>
<td>J7331</td>
<td>Hyaluronan or derivative, SYNOJOYNT, for intra-articular injection, 1 mg (new code effective 10/1/19)</td>
</tr>
<tr>
<td>J7332</td>
<td>Hyaluronan or derivative, Triluron, for intra-articular injection, 1 mg (new code effective 10/1/19)</td>
</tr>
</tbody>
</table>

**Note:** CPT codes, descriptions, and materials are copyrighted by the American Medical Association (AMA). HCPCS codes, descriptions, and materials are copyrighted by Centers for Medicare Services (CMS).
Intra-articular (IA) injection of hyaluronan into osteoarthritic joints is proposed to reduce pain and improve function. It is thought to replace endogenous hyaluronan and restore the viscoelastic properties of the synovial fluid. Most studies to date have assessed hyaluronan injections for knee osteoarthritis (OA), and this is the U.S. Food and Drug Administration-approved indication. Other joints (eg, hip, shoulder) are being investigated for IA hyaluronan treatment of OA.

Background

**Knee Osteoarthritis**

Knee osteoarthritis (OA) is common, costly, and a cause of substantial disability. Among U.S. adults, the most common causes of disability are arthritis and rheumatic disorders.

**Treatment**

Currently, no curative therapy is available for OA, and thus the overall goals of management are to reduce pain, disability, and the need for surgery.

Intra-articular (IA) injection of hyaluronan has been proposed as a means of restoring the normal viscoelasticity of the synovial fluid in patients with OA and improving pain and function. This treatment may also be called viscosupplementation. Hyaluronan is a naturally occurring macromolecule that is a major component of synovial fluid and is thought to contribute to its
viscoelastic properties. Chemical crosslinking of hyaluronan increases its molecular weight; cross-linked hyaluronans are referred to as hylans. In OA, the overall length of hyaluronan chains present in cartilage and the hyaluronan concentration in the synovial fluid are decreased.

**Summary of Evidence**

For individuals who have OA of the knee who receive IA hyaluronan injections, the evidence includes randomized controlled trials (RCTs) and systematic reviews of RCTs. The relevant outcomes are symptoms, functional outcomes, and treatment-related morbidity. Many RCTs have been published over the last 2 decades. While outcomes of these RCTs are mixed, the RCT evidence base is characterized by studies showing small treatment effects of IA hyaluronan injections. In many cases, these trials are at risk of bias, and it cannot be determined with certainty whether there is a true treatment effect or whether the reported differences are due to bias. Meta-analyses of RCTs have also had mixed findings. Some meta-analyses estimating the magnitude of treatment benefit have concluded that there is no clinically significant benefit; others have concluded that there is a clinically significant benefit. These meta-analyses have also highlighted the limitations of this evidence base, most notably publication bias and small trial bias. For example, a meta-analysis (2016) found more than a 3-fold larger treatment effect in small trials than in larger trials (ie, more than 100 participants). Overall, given the lack of a definitive treatment benefit despite a large quantity of literature, and given the biases present in the available evidence, it is unlikely that there is a treatment benefit which is clinically meaningful. The evidence is sufficient to determine that the technology is unlikely to improve the net health outcome.

For individuals who have OA of joints other than the knee who receive IA hyaluronan injections, the evidence includes RCTs, systematic reviews of RCTs, and observational studies. The relevant outcomes are symptoms, functional outcomes, and treatment-related morbidity. Meta-analyses of RCTs either have not found statistically significant benefits on health outcomes or have found benefits that were statistically, but likely not clinically, significant (eg, 0.27-point improvement on a 10-point analogue scale for hip OA). The evidence is insufficient to determine the effects of the technology on health outcomes.

**Ongoing and Unpublished Clinical Trials**

Some currently unpublished trials that might influence this review are listed in Table 1.
Table 1. Summary of Key Trials

<table>
<thead>
<tr>
<th>NCT No.</th>
<th>Trial Name</th>
<th>Planned Enrollment</th>
<th>Completion Date</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ongoing</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NCT02776514</td>
<td>Intraarticular Injections of Steroids, Hyaluronic Acid or Platelet Rich Plasma Versus Placebo for the Knee Osteoarthritis</td>
<td>240</td>
<td>Feb 2020</td>
</tr>
<tr>
<td>NCT03200288</td>
<td>A Phase 3, Randomized, Double-blind, Placebo-controlled Study to Evaluate the Clinical Performance and Safety of an Intra-articular Solution of High and Low Molecular Weight Hyaluronic Acid (HL-01) in the Treatment of Symptomatic Knee Osteoarthritis</td>
<td>691</td>
<td>Oct 2018</td>
</tr>
<tr>
<td>NCT03281837</td>
<td>A Post-market, Single Blind, Multicenter, Randomized, Controlled Trial of HYMOVIS® Intra-articular Injections in Active Subjects With Knee Osteoarthritis</td>
<td>146</td>
<td>Dec 2019</td>
</tr>
<tr>
<td><strong>Unpublished</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NCT02629380</td>
<td>Early Viscosupplementation After Partial Meniscectomy: a Double Blind, Placebo Controlled Randomized Trial</td>
<td>90</td>
<td>May 2016 (completed)</td>
</tr>
<tr>
<td>NCT02280538</td>
<td>Trial to Assess the Structural Effect and Long-term Symptomatic Relief of Intra-articular Injections of Hyaluronic Acid in Primary Knee OA (ViscOA)</td>
<td>300</td>
<td>Jan 2018 (unknown)</td>
</tr>
</tbody>
</table>

NCT: national clinical trial

* Denotes industry-sponsored or cosponsored trial

Clinical Input Received From Physician Specialty Societies and Academic Medical Centers

While the various physician specialty societies and academic medical centers may collaborate with and make recommendations during this process, through the provision of appropriate reviewers, the input does not represent an endorsement or position statement by the physician specialty societies or academic medical centers, unless otherwise noted.

In response to requests, input was received from 5 academic medical centers (6 reviewers) and 3 physician specialty societies while this policy was under review in 2011. Most reviewers agreed that intra-articular (IA) hyaluronan of the knee was medically necessary. In addition, those
providing input supported an interval of 6 months for repeat injections. In response to a question about total number of treatment courses, there was no consensus.

**Practice Guidelines and Position Statements**

**American Medical Society for Sport Medicine**

The scientific statement from the American Medical Society for Sport Medicine (2016) recommended (IA) hyaluronan for “appropriate” patients with knee osteoarthritis (OA) based on high-quality evidence. Patient selection criteria include individuals age 60 and older with Kellgren-Lawrence grade 2 or 3 OA. The society also “suggests” IA hyaluronan for patients under age 60 with knee OA based on moderate-quality indirect evidence.

**American Academy of Orthopaedic Surgeons**

The guidelines from the American Academy of Orthopaedic Surgeons (AAOS) (2013) on treatment of OA of the knee indicated that AAOS could not recommend using IA hyaluronan for patients with symptomatic knee OA. This recommendation was strong, meaning that the quality of the supporting evidence was high. It was based on a meta-analysis of 3 high-strength and 11 moderate-strength studies that showed that the overall effect was less than 0.5 minimally important different units, indicating a low likelihood that an appreciable number of patients achieved clinically important benefits. The AAOS indicated that practitioners should follow a strong recommendation unless a clear and compelling rationale for an alternative approach is present. These guidelines replaced 2008 guidelines, which included a statement that a recommendation could not be made for IA hyaluronan due to inconclusive evidence.

The AAOS (2017) clinical practice guidelines on hip OA included a recommendation that IA hyaluronic acid could not be recommended in patients with symptomatic hip OA, because it was not better than placebo. This was based on strong evidence as assessed in eight high-quality studies that evaluated IA hyaluronan against corticosteroids and placebo. Several studies showed no difference in patient pain and function after treatment with IA hyaluronan against placebo. Studies reviewing different formulations of IA hyaluronan were also considered. The AAOS (2009; reaffirmed 2014) clinical practice guidelines on glenohumeral joint OA included a weak grade C recommendation that, “The use of injectable viscosupplementation is an option when treating patients with glenohumeral [shoulder] osteoarthritis.” Grade C recommendations are based on poor-quality evidence. In this instance, the recommendation
was based on a single case series of 30 patients with OA of the glenohumeral joint who received 3 weekly IA injections of hylan G-F 20 (Synvisc). At 1, 3, and 6 months, clinically significant improvements were seen in pain, function, and quality of life measures.

**American College of Rheumatology**

The American College of Rheumatology (2012) updated is guidelines on OA of the hand, hip, and knee. A conditional recommendation was given for the use of IA hyaluronan to treat OA of the knee. The College recommended not using IA hyaluronan for OA of the hand. For OA of the hip, the College explicitly made no recommendation due to the lack of randomized controlled trials. An update is anticipated in Spring 2019.

**Osteoarthritis Research Society International**

The Osteoarthritis Research Society International (2014) guidelines, developed by consensus after review of existing guidelines and systematic reviews, gave an “uncertain” recommendation for the use of intra-articular injection of IA hyaluronan for knee OA and a recommendation of “not appropriate” for multiple-joint OA.

**National Institute for Health and Clinical Excellence**

The guidance by the National Institute for Health and Clinical Excellence (2014) stated: “Do not offer intra-articular hyaluronan injections for the management of osteoarthritis.”

**Medicare National Coverage**

There is no national coverage determination.

**Regulatory Status**

Several preparations of IA hyaluronan have been approved by the U.S. Food and Drug Administration (FDA) as an alternative to nonsteroidal anti-inflammatory drug therapy in the treatment of OA of the knee:
• Synvisc® and Synvisc-One® (Genzyme)
• Gel-One® (Zimmer)
• Hyalgan® (Fidia)
• Supartz FX™ (Bioventus)
• Orthovisc® (Anika)
• Euflexxa®, previously named Nuflexxa (Savient)
• Monovisc® (Anika Therapeutics)
• Durolane® (Bioventus)
• Gel-Syn™ (Institut Biochimique SA)

All products are manufactured from rooster combs except for Durolane®, Euflexxa®, Orthovisc®, Monovisc®, Gel-Syn™ and GenVisc® 850, which are produced from bacterial fermentation. Also, Synvisc® undergoes additional chemical crosslinking to create hylans with increased molecular weight (6000 kDa) compared with Hyalgan® (500-730 kDa) and Supartz™ (620-1170 kDa). Monovisc® is also cross-linked with a proprietary cross-linker. The differing molecular weights of the products lead to different half-lives; the half-life of Hyalgan® or Supartz™ is estimated at 24 hours, while the half-life of Synvisc® may range up to several days.

According to manufacturers’ prescribing information for Synvisc® and Euflexxa®, IA hyaluronan is “indicated for the treatment of pain in OA of the knee in patients who have failed to respond adequately to conservative nonpharmacologic therapy, and to simple analgesics, eg, acetaminophen.” The product inserts further indicate that Synvisc® and Euflexxa® should be injected intra-articularly into the knee joint once per week for a total of 3 injections over a 2- to 3-week period. In contrast, 5 weekly injections are recommended for the Hyalgan® and Supartz™ products, and 3 to 4 weekly injections are recommended for OrthoVisc®. In 2009, the FDA approved the use of single-dose hylan G-F 20 (Synvisc-One®) for the treatment of OA of the knee. In 2011, the FDA approved the use of the single-dose cross-linked hyaluronate Gel-One® (also known as Gel-200) for the treatment of OA of the knee. In 2014, Monovisc® was also approved as a single-dose treatment, while Gel-Syn™ was approved as a course of 3 weekly injections. In 2015, GenVisc 850 was approved as a course of 3 weekly injections. In 2017, Durolane was approved as a single-dose treatment.
In 2000, FDA approved removal of a precautionary statement from the package inserts for Hyalgan® and Synvisc that stated that the safety and efficacy of repeat courses have not been established.

FDA has not approved IA hyaluronan for joints other than the knee.

FDA product code: MOZ

References


History

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>05/08/12</td>
<td>New policy, add to Medicine section. This policy replaces 5.01.506. Added the table listing FDA approved hyaluranon products and recommended course(s) of treatment per MPC request. The 3rd sentence of the 4th paragraph regarding The product inserts further indicates...was moved from the description to the policy guidelines section for ease of policy administration. Policy approved with 90-day hold for provider notification. The policy effective date is November 7, 2012.</td>
</tr>
<tr>
<td>08/03/12</td>
<td>Correct Error on Related Policy number. Changed from 7.01.118 to 7.01.117.</td>
</tr>
<tr>
<td>08/27/12</td>
<td>Update Coding Section – ICD-10 codes are now effective 10/01/2014.</td>
</tr>
<tr>
<td>11/07/12</td>
<td>Policy effective after hold for provider notification. 5.01.506 is deleted.</td>
</tr>
<tr>
<td>03/08/13</td>
<td>Policy updated. Rationale and references revised. “in the knee when the above criteria are not met, and” was added to the investigational statement.</td>
</tr>
<tr>
<td>07/16/13</td>
<td>Update Related Policies. Add 7.01.549, and remove 7.01.188 as it was archived.</td>
</tr>
<tr>
<td>11/11/13</td>
<td>Replace Policy. Policy updated with literature review through July 31, 2013; reference 7 added; policy changed to not medically necessary based on new guidelines from the American Academy of Orthopaedic Surgeons. Policy change aligns with UM initiative and recent change in coverage from BCBSA. Policy held for provider notification; the effective date is April 1, 2014.</td>
</tr>
</tbody>
</table>
| 12/09/13   | Minor update: When this policy was approved last month a sentence was left out. A second policy statement is now included: “Intra-articular hyaluronan injections are
<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>01/23/14</td>
<td>Policy implementation delayed; the effective date of the policy is moved to June 1, 2014.</td>
</tr>
<tr>
<td>03/03/14</td>
<td>Policy implementation delayed; the effective date of the policy is moved to July 1, 2014.</td>
</tr>
<tr>
<td>05/09/14</td>
<td>Revised implementation date September 1, 2014. Delayed due to Plan internal system updates.</td>
</tr>
<tr>
<td>07/24/14</td>
<td>Update Related Policies. Change title to 7.01.549.</td>
</tr>
<tr>
<td>08/25/14</td>
<td>Policy implementation delayed until December 1, 2014.</td>
</tr>
<tr>
<td>12/08/14</td>
<td>Interim review. Policy updated with literature review; the policy statement is unchanged. Intra-articular hyaluronan for osteoarthritis is considered not medically necessary. Effective 03/1/15.</td>
</tr>
<tr>
<td>01/05/15</td>
<td>Coding update. New HCPCS code J7327 added to the policy.</td>
</tr>
<tr>
<td>02/09/15</td>
<td>Policy implementation date extended to May 1, 2015.</td>
</tr>
<tr>
<td>03/13/15</td>
<td>Policy implementation date extended to June 1, 2015.</td>
</tr>
<tr>
<td>03/24/15</td>
<td>Update Related Policies. Change title to 7.01.549.</td>
</tr>
<tr>
<td>04/14/15</td>
<td>Implementation update: Policy will now be effective July 1, 2015. In the interim, see policy 2.01.534 (effective 4/14/15) for coverage (link provided in header).</td>
</tr>
<tr>
<td>06/01/15</td>
<td>Coding update. ICD-9 codes removed; these are not utilized in policy adjudication.</td>
</tr>
<tr>
<td>12/08/15</td>
<td>Annual Review. Policy reviewed with literature search. No change to policy statements.</td>
</tr>
<tr>
<td>01/19/16</td>
<td>Coding update. New HCPCS codes J7328 and Q9980, effective 1/1/16, added to policy.</td>
</tr>
<tr>
<td>09/30/16</td>
<td>Policy moved into new format; no change to policy statements.</td>
</tr>
<tr>
<td>01/01/17</td>
<td>Coding update; added new HCPCS codes J7320 and J7322 effective 1/1/17.</td>
</tr>
<tr>
<td>01/01/18</td>
<td>Removed HCPCS code Q9980 as it terminated on 1/1/17.</td>
</tr>
<tr>
<td>07/01/18</td>
<td>Annual Review, approved June 5, 2018. Policy updated with literature review through February 2018; references 29 and 33 added. Policy statements unchanged.</td>
</tr>
<tr>
<td>01/01/19</td>
<td>Coding update, added new HCPCS codes J7318 and J7329 (new codes effective 1/1/19)</td>
</tr>
</tbody>
</table>
### Comments

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
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</thead>
<tbody>
<tr>
<td>10/01/19</td>
<td>Coding updated, added HCPCS J7331 and J7332 (new codes effective 10/1/19).</td>
</tr>
</tbody>
</table>

**Disclaimer:** This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. The Company adopts policies after careful review of published peer-reviewed scientific literature, national guidelines and local standards of practice. Since medical technology is constantly changing, the Company reserves the right to review and update policies as appropriate. Member contracts differ in their benefits. Always consult the member benefit booklet or contact a member service representative to determine coverage for a specific medical service or supply. CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). ©2019 Premera All Rights Reserved.

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Toll free 855-332-4535, Fax 425-918-5592, TTY 800-842-5357
Email AppealsDepartmentInquiries@Premera.com

You can file a grievance in person or by mail, fax, or email. If you need help filing a grievance, the Civil Rights Coordinator 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 https://ocrportal.hhs.gov/ocr/portal/lobby.jsf, or by mail or phone at:
U.S. Department of Health and Human Services
200 Independence Avenue SW, Room 509F, HHH Building
Washington, D.C. 20201, 1-800-368-1019, 800-537-7697 (TDD)

Getting Help in Other Languages

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Call 800-722-1471 (TTY: 800-842-5357).

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يجب أن تكون هذه المعلومات مفيدة، وفي كل حال، فإنها لم تكن مفيدة في هذا الإشعار. وقد تحتاج لإعادة ترتيب المعلومات على ترتيب يمكن أن يكون الرفرف على كيفية استعمال المعلومات المصممة باللغة العربية. في ذلك التحصيل، ي.viewmodel لو أنك تجد المعلومات على هذه المعلومات والمعلومات والاستخدام الأخرى من ال kişiler. في كل مكان:

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Lakkoofsa biliibila 800-722-1471 (TTY: 800-842-5357) ti biliibila.

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Appelez le 800-722-1471 (TTY: 800-842-5357).

Kreyol Ayisyen (Creole):

Avi sila a gen Enfòmasyon Enpòtan ladan. Avi sila a kapab genyen enfòmasyon enpòtan konsèn an aplanasyon w lan oswa konsèn kouvèti asirans lan atravè Premera Blue Cross. Kapab genyen dat ki enpòtan nan avi sila a. Ou ka gen pou pran kék aplanasyon avan sèten dat limit pou ka kenbe kouvèti asirans santre w lan oswa pou yo ka ede w avèk depans yo. Se dwa w pou resewa enfòmasyon sa a ak asistans nan lang ou pa ale a, san ou pa gen pou peye pou sa. Rate nan 800-722-1471 (TTY: 800-842-5357).

Deutsche (German):


Hmooj (Hmong):

Tsab ntawv tshaaj xo no muaj cov ntsiab lus tseem ceeb. Tej zaum tsab ntawv tshaaj xo no muaj cov ntsiab lus tseem ceeb bokj kja daim ntawv thov kev pab los yoy koj qhov kev pab cuam los ntawm Premera Blue Cross. Tej zaum muaj cov hnb tseem ceeb uss sau rau hauv daim ntawv no. Tej zaum kja yuav ta joo yam uss peb kom kja us tib puab kaj nyooy us tib tsev rau hauv daim ntawv no. Tej zaum kja yuav ta joo yam uss peb kom kja us tib puab kaj nyooy us tib tsev rau hauv daim ntawv no.
Muaj cov nuj bna tsawv nhia joo yam uss peb kom kja us tib puab kaj nyooy us tib tsev rau hauv daim ntawv no.

Illoko (Illoko):

Daytoy a Pakdaak ket naglaon iti Napateg nga Impormasion. Daytoy a pakdaak mabalin nga adda ket naglaon iti napateg nga impormasion maiupayeep iti aplikasyonu yowo coverage babaen iti Premera Blue Cross. Daytoy kaj mabalin dagiti importante a pelsa iti daytoy a pakdaak. Mabalin nga adda rumbeng nga aramidenyo nga addag sakbay dagiti partikular a naituing nga adda algaw tapno mapagalatneddyo ti coverage ti salun-ti yowo tongad tagadiy gastos. Adda karbenganyo a manga iti daytoy nga impormasion ken tongul iti bukodyo a pasasasao nga awan ti bayaadanyo. Tumawag ditu numero nga 800-722-1471 (TTY: 800-842-5357).

Italiano (Italian):

Questo avviso contiene informazioni importanti. Questo avviso può contenere informazioni importanti sulla tua domanda o copertura attraverso Premera Blue Cross. Potrebbero esserci date chiave in questo avviso. Potrebbe essere necessario un tuo intervento entro una scadenza determinata per consentirti di mantenere la tua copertura o sovvenzione. Hai il diritto di ottenere queste informazioni e assistenza nella tua lingua gratuitamente.
Chiama 800-722-1471 (TTY: 800-842-5357).

Latin America: This Notice contains Important Information. This notice may have important information about your application or coverage through Premera Blue Cross. There may be key dates in this notice. You may need to take action before a deadline to keep your health coverage or benefits. You have the right to get this information and help in your language at no cost.
Call 800-722-1471 (TTY: 800-842-5357).

Spanish: Este aviso contiene información importante. Este aviso puede contener información importante sobre tu solicitud o cobertura a través de Premera Blue Cross. Pueden haber fechas clave en este aviso. Podrías necesitar actuar antes de una fecha límite para mantener tu cubrimiento de salud o ayudas. Tienes derecho a obtener esta información y ayuda en tu idioma gratuitamente.
Português (Portuguese):
Este aviso contém informações importantes. Este aviso poderá conter informações importantes a respeito de sua aplicação ou cobertura por meio do Premera Blue Cross. Poderão existir datas importantes neste aviso. Talvez seja necessário que você tome providências dentro de determinados prazos para manter sua cobertura de saúde ou ajuda de custos. Você tem o direito de obter esta informação e ajuda em seu idioma e sem custos. Ligue para 800-722-1471 (TTY: 800-842-5357).

Română (Romanian):

Русский (Russian):
Настоящее уведомление содержит важную информацию. Это уведомление может содержать важную информацию о вашем заявлении или страховом покрытии через Premera Blue Cross. В настоящем уведомлении могут быть ключевые даты. Вам, возможно, потребуется принять меры к определенным предельным срокам для сохранения страхового покрытия или помощи с расходами. Вы имеете право на бесплатное получение этой информации и помощь на вашем языке. Звоните по телефону 800-722-1471 (TTY: 800-842-5357).

Español (Spanish):
Este Aviso contiene información importante. Es posible que este aviso contenga información importante acerca de su solicitud o cobertura a través de Premera Blue Cross. Es posible que haya fechas clave en este aviso. Es posible que deba tomar alguna medida antes de determinadas fechas para mantener su cobertura médica o ayuda con los costos. Usted tiene derecho a recibir esta información y ayuda en su idioma sin costo alguno. Líame al 800-722-1471 (TTY: 800-842-5357).

Romania (Romanian):

Polski (Polish):

Tiếng Việt (Vietnamese):

Український (Ukrainian):
Це повідомлення містить важливу інформацію. Це повідомлення може містити важливу інформацію про Ваше звернення щодо страхувального покриття через Premera Blue Cross. Зверніть увагу на ключові дати, які можуть бути вказані у цьому повідомленні. Існує імовірність того, що Вам треба буде здійснити певні кроки у конкретні кінцеві строки для того, щоб зберегти Ваше медичне страхування або отримати фінансову допомогу. У Вас є право на отримання цієї інформації та допомоги безкоштовно на Вашій рідній мові. Дозвоніться за номером телефону 800-722-1471 (TTY: 800-842-5357).

日本語 (Japanese):
この通報に重要な情報が含まれています。この通報、Premera Blue Crossの申請または補償範囲に関する重要な情報が含まれています。この通報に記載されている情報が重要な日常生活をご確認ください。健康保険や無料サポートを維持するには、特定の期限までに行動を取られなければなりません。ご用件の言語による情報とサポートが無料で提供されます。800-722-1471 (TTY: 800-842-5357) までお電話ください。

한국어 (Korean):
본 통지서에는 중요한 정보가 들어 있습니다. 즉 이 통지서는 귀하의 신청에 관하여 그리고 Premera Blue Cross를 통한 커버리지에 관한 정보를 포함하고 있을 수 있습니다. 귀하의 신청과 커버리지를 계속 유지하거나 비용을 절감하기 위해서 일정한 마감일까지 조치를 취해야 할 필요가 있을 수 있습니다. 귀하의 이런 정보와 정보를 귀하의 안내로 비용 부담이 없을 수 있는 권리가 있습니다. 800-722-1471 (TTY: 800-842-5357) 로 전화하시오.