

San Francisco Health Plan

Healthy Workers HMO

Formulario de medicamentos recetados

A partir del August 2025

Este formulario se actualizó el <Date of Printable formulary posting in August 20,2025. Este formulario está sujeto a cambios, y todas las versiones anteriores del formulario ya no están en vigor.

Formulario más reciente:

sfhp.org/for-members/healthy-workers/benefits/pharmacy-services

Beneficios y servicios de HW:

sfhp.org/programs/healthy-workers/benefits

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Sección informativa

Introducción al Formulario

El Formulario de San Francisco Health Plan (SFHP) es una lista de productos farmacéuticos aprobados por la Food and Drug Administration (FDA) y elegibles para la cobertura del beneficio de medicamentos recetados para pacientes ambulatorios. Los medicamentos que se incluyen en el Formulario de SFHP se seleccionan para reflejar las terapias farmacológicas más apropiadas, de alta calidad y rentables. El Formulario SFHP incluye medicamentos de marca y genéricos, así como algunos dispositivos, de conformidad con la Affordable Care Act (ACA) y las regulaciones estatales de California. La presencia de un medicamento en el Formulario no garantiza que su proveedor le recetará ese medicamento.

Definición de términos

La siguiente es una lista de términos comunes utilizados en este Formulario y sus significados:

"Medicamento de marca" es un medicamento que se comercializa bajo un nombre de marca registrada protegida. El medicamento de marca debe aparecer en letras MAYÚSCULAS.

"Coseguro" es un porcentaje del costo de un beneficio de atención médica cubierto que un afiliado paga después de que el afiliado ha pagado el deducible, si se aplica un deducible al beneficio de atención médica, como el beneficio de medicamentos con receta.

"Copago" es un monto fijo en dólares que un afiliado paga por un beneficio de atención médica cubierto después de que el afiliado ha pagado el deducible, si se aplica un deducible al beneficio de atención médica, como el beneficio de medicamentos recetados.

"Deducible" es el monto que un afiliado paga por los beneficios de atención médica cubiertos antes de que el plan de salud del afiliado comience a pagar la totalidad o parte del costo del beneficio de atención médica conforme a los términos de la póliza.

"Nivel del medicamento" se refiere a un grupo de medicamentos recetados que corresponde a un nivel de costo compartido especificado en la cobertura de medicamentos recetados del plan de salud. El nivel en el que se coloca un medicamento recetado determina la parte del costo del medicamento para el afiliado.

"Afiliado o persona inscrita" es la persona que es miembro de un plan de salud y recibe servicios a través del plan. Todas las referencias a los afiliados en esta plantilla

de formulario también incluirán al suscriptor, tal como se define en esta sección a continuación.

“Solicitud de excepción” es una solicitud de cobertura de un medicamento recetado. Si un afiliado, su designado o el proveedor de atención médica que receta presenta una solicitud de excepción para la cobertura de un medicamento recetado, el plan de salud debe cubrir el medicamento recetado cuando se determine que es médica mente necesario para tratar la afección del afiliado.

“Circunstancias apremiantes” son aquellas que surgen cuando un afiliado tiene una condición de salud que podría poner gravemente en peligro su vida, su salud o su capacidad de recuperar su función máxima o cuando un afiliado se somete a un curso de tratamiento actual usando un medicamento que no se incluye en el formulario.

“Formulario” es la lista completa de medicamentos preferidos para uso y que cumple con los requisitos para cobertura bajo un producto del plan de salud, e incluye todos los medicamentos cubiertos bajo el beneficio de medicamentos recetados para pacientes ambulatorios del producto del plan de salud. El Formulario también se conoce como Lista de medicamentos recetados.

“Medicamento genérico” es el mismo medicamento que su equivalente de marca en cuanto a dosis, seguridad, potencia, modo de administración, calidad, rendimiento y uso previsto. Un medicamento genérico figura en ***negrita y en minúscula cursiva***.

“Medicamento no incluido en el Formulario” es un medicamento recetado que no figura en el formulario del plan de salud.

“Costos de desembolso directo” son los copagos, el coseguro y el deducible aplicable, además de todos los costos por servicios de atención médica que no están cubiertos por el plan de salud.

“Comité de Farmacia y Terapéutica” es un grupo de personas que recetan y farmacéuticos locales que se reúne cuatro veces al año y determinan qué medicamentos se incluirán en el Formulario y los criterios usados para el proceso de revisión de Autorización previa.

“Proveedor que receta” es un proveedor de atención médica autorizado para emitir una receta para tratar una afección médica de un afiliado al plan de salud.

“Receta” es una orden oral, escrita o electrónica de un proveedor que receta para un afiliado específico que contiene el nombre del medicamento recetado, la cantidad del medicamento recetado, la fecha de emisión, el nombre y la información de contacto del proveedor que receta, la firma del proveedor que receta si la receta es por escrito y, si lo solicita el afiliado, la afección médica o el propósito para el que se receta el medicamento.

“Medicamento recetado” es un medicamento recetado por el proveedor que receta del afiliado y que requiere una receta de conformidad con la ley aplicable.

“Autorización previa” es el requisito del plan de salud para que el afiliado o el proveedor que receta al afiliado obtengan la autorización del plan de salud para un medicamento recetado antes de que el plan de salud cubra el medicamento. El plan de salud otorgará una autorización previa cuando sea médicaamente necesario que el afiliado obtenga el medicamento.

“Terapia escalonada o terapia de pasos” es un proceso que especifica la secuencia en la que se receten diferentes medicamentos recetados para una determinada afección médica y médicaamente apropiados para un paciente en particular. El plan de salud puede requerir que el afiliado pruebe uno o más medicamentos para tratar la afección médica del afiliado antes de que el plan de salud cubra un medicamento en particular para la afección conforme a una solicitud de terapia escalonada. Si el proveedor que receta del afiliado presenta una solicitud de excepción de terapia escalonada, los planes de salud harán excepciones a la terapia escalonada cuando se cumplan los criterios.

“Suscriptor” se refiere a la persona que es responsable del pago a un plan o cuyo empleo u otra condición, excepto por dependencia familiar, es la base para la elegibilidad para la afiliación al plan.

Cómo leer este Formulario

El documento del Formulario de SFHP está ordenado alfabéticamente por clase de medicamento e incluye la siguiente información: nombre del medicamento, forma farmacéutica, nivel del medicamento, junto con cualquier restricción del Formulario como límite de cantidad, autorización previa o requisitos de terapia escalonada.

¿Cómo puedo encontrar un medicamento en la lista del Formulario?

Puede encontrar un medicamento recetado buscando la categoría terapéutica y la clase del medicamento o la MARCA o el nombre **genérico** del medicamento en el índice alfabético. Puede buscar en esta lista:

- Buscando la categoría o clase a la que pertenece el medicamento y luego buscar el nombre del medicamento en orden alfabético; O bien
- Buscando el nombre del medicamento en el índice en orden alfabético

Lista del Formulario

| Tipo de medicamento | Cómo aparecerá el nombre del medicamento en la lista de medicamentos del Formulario |
|--|---|
| medicamento genérico | metronidazol, comprimido oral |
| medicamento genérico con un nombre de marca comercializado | norgestimato (Tri-Sprintec) |
| medicamento de marca | ELIQUIS, COMPRIMIDO ORAL (apixabán) |

¿Cómo sé si el medicamento que aparece en la lista es de marca o genérico?

Un medicamento aparece ordenado alfabéticamente por su MARCA y sus nombres **genéricos** en la categoría terapéutica y clase a la que pertenece. Si un medicamento sólo está disponible como marca, el nombre genérico se incluye después del nombre de marca entre paréntesis y todo **en negrita y minúsculas cursivas**. Si un medicamento está disponible como genérico, se incluye por el nombre genérico. Algunos medicamentos genéricos se comercializan bajo una marca patentada y protegida. Para estos, el nombre de la marca aparecerá después del nombre genérico entre paréntesis y el tipo de letra regular con la primera letra de cada palabra en mayúscula.

Por lo general, SFHP requiere la sustitución por medicamento genérico cuando se dispone de un producto genérico equivalente a un medicamento de marca (consulte la Política de medicamentos de marca a continuación). Si existe disponible un equivalente genérico para un medicamento de marca y tanto el equivalente de marca como el genérico están cubiertos, el medicamento genérico aparecerá por separado del medicamento de marca **en negrita y minúsculas cursivas**. Si el equivalente genérico de un medicamento de marca no está disponible o no está cubierto en el Formulario, el medicamento no aparecerá en la lista separada por su nombre genérico.

¿Qué sucede si no puedo encontrar un medicamento en la lista del Formulario?

Si su medicamento no figura en la lista del Formulario de SFHP, se denomina medicamento no incluido en el Formulario. Su proveedor debe enviar un formulario de autorización previa a SFHP antes de que la farmacia pueda dispensar el medicamento que no está en el Formulario. SFHP revisará la solicitud y determinará si el medicamento será autorizado según los criterios de Autorización previa aprobados por el Comité de Farmacia y Terapéutica de SFHP. Algunos medicamentos del Formulario también requieren que su proveedor envíe un formulario de autorización previa antes de que la farmacia pueda dispensar el medicamento si su receta excede los límites específicos, requiere terapia escalonada o es un medicamento con usos específicos. El criterio de Autorización previa de SFHP está aprobado por el Comité de SFHP P&T y es consistente con las directrices de tratamiento reconocidas profesionalmente y estándares de práctica.

*Para obtener más información sobre el proceso de autorización previa, consulte la sección Restricciones del Formulario.

¿Qué son los Niveles de medicamentos?

Los medicamentos se colocan en diferentes niveles según su funcionamiento, su seguridad y su costo en comparación con otros medicamentos utilizados para el mismo tipo de tratamiento.

- Los medicamentos del Nivel 1 son medicamentos genéricos del Formulario. Pueden aplicarse límites de cantidad y edad.
- Los medicamentos de Nivel 2 son medicamentos de marca del formulario. Pueden aplicarse límites de cantidad y edad.
- Los medicamentos del Nivel 3 son medicamentos genéricos o de marca del Formulario que requieren una autorización previa o requieren terapia escalonada.
- Los medicamentos sin nivel son medicamentos no incluidos en el formulario, medicamentos excluidos o medicamentos cubiertos a través del beneficio médico.

| Nivel del medicamento | | Requisitos y límites de cobertura* |
|-----------------------|---|---|
| Nivel 1 | Formulario, genérico | AL = Límite de edad QL = Límite de cantidad |
| Nivel 2 | Formulario, marca | |
| Nivel 3 | Formulario con autorización previa o terapia escalonada requerida (puede ser de marca o genérico) | PA = Autorización previa ST = Terapia escalonada |

*El nivel de medicamento y otras restricciones se designan con símbolos de acuerdo con la clave de arriba. Consulte la sección “Restricciones del Formulario” para obtener más detalles sobre estos límites.

Algunos medicamentos pueden figurar en la lista en varios niveles debido a que una concentración en particular es un formulario y otra forma de concentración o dosificación del mismo medicamento requiere una autorización previa.

¿Cuál es la diferencia entre el beneficio de medicamentos recetados para pacientes ambulatorios y el beneficio médico?

El beneficio de medicamentos recetados para pacientes ambulatorios incluye medicamentos aprobados por la FDA que son autoadministrados, comúnmente medicamentos orales o autoinyectables, no excluidos de otra manera. Los medicamentos que deben ser administrados por un profesional de la atención médica generalmente se proporcionan bajo el beneficio médico. Los medicamentos del beneficio médico pueden incluir medicamentos para terapia de infusión, hemoderivados y cualquier fármaco que deba ser administrado por un profesional de atención médica. Los medicamentos del beneficio médico no están cubiertos en el beneficio de medicamentos recetados para pacientes ambulatorios, con las siguientes excepciones: medicamentos inyectables de acción prolongada para el tratamiento necesario de una

afección de salud mental o trastorno por consumo de sustancias, y un medicamento inyectable utilizado para la profilaxis previa a la exposición (PrEP) para el HIV-1. Para obtener la lista completa de medicamentos que debe administrarle un profesional de atención médica que también están cubiertos por el beneficio de medicamentos recetados para pacientes ambulatorios, e información sobre cómo obtener medicamentos a través del beneficio médico, visite sfhp.org/for-members/healthy-workers/benefits/pharmacy-services

Para obtener más información sobre los beneficios del plan Healthy Workers HMO, visite sfhp.org/programs/healthy-workers/benefits.

¿Hay algún medicamento excluido del Formulario?

Las siguientes clases de medicamentos están excluidas del Formulario de Healthy Workers HMO:

- Medicamentos para el uso de la disfunción erétil, excepto cuando se recetan como un tratamiento médica mente necesario de una afección de salud mental o trastorno por uso de sustancias.
- Productos de medicamentos compuestos cuando están aprobados por la FDA y productos comercializados disponibles para el diagnóstico. También se debe haber comprobado la seguridad, eficacia y estabilidad de estos productos compuestos para la consideración de una excepción para esta exclusión.
- Medicamentos administrados por un profesional de atención médica, excepto medicamentos inyectables de acción prolongada para el tratamiento necesario de una afección de salud mental o trastorno por consumo de sustancias y un medicamento inyectable utilizado para la profilaxis previa a la exposición (PrEP) para el HIV-1.
- Vitaminas de venta libre (OTC) que no sean médica mente necesarias, medicamentos OTC y dispositivos OTC. Las excepciones incluyen: aspirina para prevenir enfermedades cardiovasculares y cáncer colorrectal para adultos de 50–59 años con un alto riesgo cardiovascular, suministros para diabéticos, dispositivos anticonceptivos y medicamentos, suministros y dispositivos para el tratamiento de la fenilcetonuria (PKU), tratamiento médica mente necesario de una afección de salud mental o trastorno de salud mental o por consumo de sustancias, medicamentos para ayudarle a dejar de fumar y vitaminas prenatales, que incluyen preparaciones de ácido fólico y flúor si son médica mente necesarios y requieren una receta médica.
- Las recetas de medicamentos o dispositivos que no han recibido la aprobación por parte de la FDA están excluidos.

¿Quién decide qué medicamentos están incluidos en el Formulario?

El Comité de Farmacia y Terapéutica de SFHP (P&T) es responsable de seleccionar los medicamentos en el Formulario. El Comité de SFHP P&T está compuesto por médicos de la red que participan activamente de diversas especialidades y clínicas médicas, así como por farmacéuticos clínicos comunitarios junto con el director médico y director de farmacia de SFHP o su designado. El Comité de SFHP P&T revisa todos los medicamentos nuevos y las nuevas pautas de tratamiento para determinar qué tan bien funcionan los medicamentos, el perfil de seguridad de los medicamentos y el valor general al seleccionar la lista del formulario.

Los proveedores de SFHP pueden solicitar la evaluación de medicamentos para agregarlos o eliminarlos del formulario enviando el Formulario de solicitud de modificación al formulario disponible en nuestro sitio web en sfhp.org/providers/pharmacy-services/prior-authorization-requests.

¿Cómo y cuándo cambia el Formulario?

El Comité de P&T se reúne trimestralmente en enero, abril, julio y octubre para revisar los cambios al Formulario en función de consideraciones de seguridad, eficacia y calidad de la atención. Las actualizaciones trimestrales del Formulario aprobadas durante las reuniones del Comité de P&T entran en vigor el día 20 del mes siguiente y se publican en el sitio web de SFHP para su revisión. Otros cambios o actualizaciones provisionales también se publicarán mensualmente para su revisión. Una vez actualizados, los documentos anteriores del Formulario ya no se consideran en vigor.

Pueden ocurrir los siguientes cambios en el Formulario:

- Un medicamento puede agregarse o retirarse del formulario
- Se puede agregar un nuevo formulario genérico al formulario cuando esté disponible
- Un medicamento de marca puede ser retirado del Formulario cuando esté disponible un genérico igual
- Un medicamento puede cambiar de nivel cuando se agregan o eliminan requisitos de autorización previa o de terapia escalonada
- Se pueden agregar, eliminar o cambiar los límites de edad o cantidad

Cuando un medicamento o forma de dosificación se retire del Formulario o se agreguen restricciones y haya sido aprobado previamente para la cobertura de su afección médica, la cobertura del medicamento continuará si su proveedor continúa recetándole el medicamento para su afección y el medicamento se receta de manera adecuada y es seguro para su afección. SFHP le notificará si un medicamento cubierto que está tomando se retira del formulario porque la FDA considera que el medicamento no es seguro y se retira del mercado, o porque el fabricante lo retira del mercado.

Puede acceder a las actualizaciones mensuales del Formulario de SFHP en línea desde nuestro sitio web en sfhp.org/for-members/healthy-workers/benefits/pharmacy-services. También puede solicitar información llamando a Servicio al Cliente de SFHP al **1(415) 547-7800** (local), **1(800) 288-5555** (llamada gratuita), de lunes a viernes, de 8:30am–5:30pm.

¿Qué servicios preventivos están cubiertos por el beneficio de medicamentos recetados para pacientes ambulatorios?

La Patient Protection and Affordable Care Act (ACA) y la Knox-Keene Health Care Service Plan Act (Knox-Keene Act) exigen que los servicios preventivos estén cubiertos sin autorización previa y sin cargo para el afiliado. Los medicamentos de salud preventiva cubiertos por el beneficio de medicamentos recetados para pacientes ambulatorios se determinan con base en las recomendaciones del United States Preventive Services Task Force y la cobertura de vacunas se basa en las recomendaciones del Advisory Committee on Immunization Practices de los Centers for Disease Control and Prevention federales. Para obtener más detalles sobre los servicios preventivos cubiertos por el beneficio de medicamentos recetados para pacientes ambulatorios, visite sfhp.org/for-members/healthy-workers/benefits/pharmacy-services. Para obtener información sobre cómo obtener medicamentos recetados y cómo encontrar una farmacia de la red, consulte la sección “Cómo surtir una receta” a continuación.

¿Qué es un medicamento o dispositivo anticonceptivo?

Los anticonceptivos son medicamentos o dispositivos, como los diafragmas, que ayudan a prevenir el embarazo. SFHP está obligado por ley a cubrir un suministro para hasta 12 meses, por surtido, de medicamentos y dispositivos anticonceptivos aprobados por la FDA, incluidos los anticonceptivos de venta libre (OTC), sin autorización previa y sin copago. No se requiere receta médica para activar la cobertura de los anticonceptivos OTC. La tarjeta de identificación del afiliado debe presentarse en una farmacia participante dentro de la red y la farmacia puede procesar el anticonceptivo OTC del formulario sin autorización previa y sin copago.

Los anticonceptivos de venta libre (OTC) incluyen:

- Condones (femenino)
- Condones (masculino)
- Anticonceptivos orales diarios (Opill)
- Anticonceptivos orales de emergencia
- Espermicidas (crema, película, espuma, gel, suppositorio)

Para obtener información sobre cómo obtener un anticonceptivo recetado y cómo encontrar una farmacia de la red, consulte la sección “Cómo surtir una receta” a continuación.

¿Qué medicamentos y productos para el cuidado de la diabetes están cubiertos?

SFHP cubre los medicamentos aprobados por la FDA para el tratamiento de la diabetes (incluido el tipo de diabetes que se desarrolló durante el embarazo en mujeres que no habían tenido diabetes anteriormente) y los dispositivos y suministros para las pruebas de diabetes, ya sea que necesite o no insulina. Los dispositivos y suministros cubiertos por el beneficio de medicamentos recetados para pacientes ambulatorios incluyen monitores de glucosa, tiras reactivas, jeringas, lancetas, toallitas impregnadas en alcohol y tiras reactivas de cetonas en orina. Para obtener información sobre cómo obtener medicamentos recetados y cómo encontrar una farmacia de la red, consulte “Cómo surtir una receta” a continuación.

Cómo surtir una receta

Cuando necesite medicamentos, su médico de atención primaria (PCP) o el especialista al cual fue derivado se los recetará. Puede obtener medicamentos recetados en cualquier farmacia minorista dentro de la red. Para obtener el medicamento, lleve la receta impresa a un farmacia minorista que figure en la sección Farmacias del Directorio de proveedores de San Francisco Health Plan Healthy Workers HMO (sfhp.org/programs/healthy-workers/find-a-provider) y muestre su tarjeta de identificación de afiliado al farmacéutico. Su médico de atención primaria (PCP) o especialista al que fue derivado puede optar por enviar los medicamentos recetados a una farmacia de forma electrónica. Las recetas se cubrirán de acuerdo con este documento del Formulario de SFHP y las restricciones que se describen a continuación.

Los afiliados también pueden obtener sus medicamentos a través de una farmacia de pedidos por correo. Puede ser una forma conveniente de surtir medicamentos de mantenimiento. Los medicamentos de mantenimiento son fármacos que los médicos receten de forma continua y regular para mantener la salud. Para ver instrucciones sobre cómo obtener medicamentos a través de una farmacia de pedidos por correo, visite la sección de Farmacia de pedidos por correo en el siguiente sitio web sfhp.org/programs/healthy-workers/benefits/pharmacy-services.

Los medicamentos que figuran en el formulario de SFHP Healthy Workers HMO no están sujetos a ninguna restricción de farmacia especializada ni a ninguna otra limitación de cobertura de la red. SFHP Healthy Workers HMO no cuenta con farmacias especializadas en la red.

Información sobre el copago:

Los copagos por medicamentos recetados cubiertos para pacientes ambulatorios son los que se describen a continuación. El costo compartido no excede el 50 por ciento del costo para el plan.

Los siguientes copagos se aplican a las recetas cubiertas por el Formulario de Healthy Workers HMO:

- **Copago de \$5:** medicamentos genéricos incluidos en el Nivel 1 o 3, y medicamentos de marca preferidos incluidos en el Nivel 2 o 3 que tienen un equivalente genérico.
- **Copago de \$10:** medicamentos de marca incluidos en el Nivel 2 o Nivel 3

El monto total de copagos y coseguro que debe pagar un afiliado no excede los doscientos cincuenta dólares (\$250) por un suministro para hasta 30 días de un medicamento recetado cubierto para pacientes ambulatorios.

¿A qué medicamentos se les exime el costo?

La legislación estatal o federal exige que algunos medicamentos estén cubiertos, sin que los afiliados deban asumir ningún desembolso directo. Para ver la lista completa de medicamentos preventivos y vacunas a \$0 de copago, visite

sfhp.org/programs/healthy-workers/benefits/pharmacy-services.

- **Copago de \$0:**
 - medicamentos preventivos, que incluyen, entre otros:
 - todos los anticonceptivos
 - medicamentos aprobados para la prevención de la infección por el virus de inmunodeficiencia humana (HIV)
 - aspirina de dosis baja
 - medicamentos con estatinas
 - vitaminas prenatales con ácido fólico
 - medicamentos para la prevención del cáncer de mama
 - medicamentos para dejar de fumar
 - vacunas
 - Pruebas caseras y tratamientos para el Covid-19 proporcionados en las farmacias de la red de SFHP.
- Todos los medicamentos con copago de \$0 en el Formulario aparecen con este símbolo: 

Restricciones del Formulario

El Formulario de SFHP utiliza restricciones estándar del formulario designadas con símbolos que incluyen límites específicos para los medicamentos, como límites de cantidad (QL) y límites de edad (AL), terapia de pasos (ST) y autorización previa (PA). Todas las restricciones del formulario se basan en indicaciones, normas de práctica y consideraciones de seguridad aprobadas por la FDA. Las recetas que excedan las restricciones del Formulario requieren que su profesional que receta envíe una solicitud de autorización previa. Su farmacéutico recibirá un mensaje electrónico de SFHP si una receta tiene una restricción del Formulario que requiere autorización previa y le informará a usted y a su proveedor tratante.

¿Qué es la terapia escalonada?

Los medicamentos de terapia escalonada (ST) requieren que pruebe uno o más medicamentos para tratar su afección médica antes de que SFHP cubra un medicamento en particular para la afección de conformidad con una solicitud de terapia escalonada. Si el proveedor que le receta presenta una solicitud de excepción de terapia escalonada, SFHP hará excepciones a la terapia escalonada cuando se cumpla con los criterios de autorización previa. SFHP no exigirá que cumpla con una terapia escalonada para un medicamento que ya esté tomando, siempre y cuando su proveedor continúe recetándole el medicamento y el medicamento siga siendo apropiado y se considere seguro y eficaz para su afección. Para solicitar una exención, se debe presentar una autorización previa utilizando los [criterios de excepción de terapia escalonada](#). Puede encontrar información adicional para la aprobación de la solicitud de terapia escalonada en sfhp.org/providers/pharmacy-services/prior-authorization-requests bajo la sección de “Criterio de autorización previa”.

¿Cuál es el proceso de solicitud de autorización previa?

El plan de salud cubrirá medicamentos que no estén incluidos en el formulario o medicamentos restringidos cuando sean médica mente necesarios. Si un medicamento que no está en el Formulario, un medicamento restringido para autorización previa, un medicamento restringido para terapia escalonada o una receta que excede un límite de cantidad o edad es médica mente necesario; usted o su proveedor pueden solicitar a SFHP que revise la receta para la cobertura. Este proceso se denomina solicitud de autorización previa o solicitud de excepción. Los médicos clínicos pueden enviar una solicitud de autorización previa de la siguiente manera:

1. **Por fax:** Descargue un [Formulario de solicitud de autorización previa](#) y envíelo por fax al **1(855) 461-2778** tanto para las solicitudes estándar como las urgentes. Las solicitudes urgentes deben estar claramente etiquetadas como “URGENTE” en la parte superior del formulario de solicitud de autorización previa.
2. **Por teléfono:** Llame al administrador de beneficios de farmacia (PBM) Magellan al **1(800) 424-4331** para presentar una solicitud oral.

Puede acceder al [Formulario de solicitud de autorización previa](#) desde nuestro sitio web en sfhp.org/providers/pharmacy-services/prior-authorization-requests.

Un farmacéutico o director médico revisa todas las solicitudes de autorización previa y toma la decisión de aprobar, aprobar con cambios, denegar o pedirle más información al proveedor que le receta con base en los criterios aprobados por el Comité de SFHP P&T. Las solicitudes no urgentes se revisan en un plazo de 72 horas. Cuando existan circunstancias apremiantes, la solicitud se procesa como expedita y se revisa en 24 horas. Cuando existan circunstancias apremiantes, la solicitud puede etiquetarse como urgente. Si la solicitud de autorización previa se aprueba, se envía un mensaje por fax a la persona que receta que envió el Formulario de solicitud de autorización previa indicando que SFHP cubrirá el medicamento. Cuando se aprueba una solicitud que no es urgente, el plan de salud proporcionará cobertura durante el tiempo que dure la receta, incluyendo los resurtidos. Cuando se aprueba una solicitud basada en circunstancias apremiantes, el plan de salud proporcionará cobertura durante el tiempo que dure la exigencia. Si el plan de salud no responde a una solicitud completa de autorización previa o de terapia escalonada dentro de las 72 horas posteriores a la recepción de una solicitud no urgente o 24 horas posteriores a la recepción de una solicitud basada en circunstancias apremiantes, la solicitud se considerará concedida por el tiempo que dure la receta, incluidos los resurtidos.

Si se deniega o cambia la autorización previa, SFHP les enviará una carta a usted y al proveedor que receta. Esta carta incluye el motivo de la decisión de SFHP. También incluimos instrucciones sobre cómo puede apelar si no está de acuerdo con la decisión. Si no está de acuerdo con la denegación o aprobación con modificaciones de SFHP, puede presentar una apelación ante SFHP. SFHP revisará su apelación en un plazo de treinta (30) días. Si su apelación es urgente, se revisará dentro de 72 horas. Si su apelación es acerca de un medicamento que no está en el formulario de SFHP, puede presentar una queja para pedir una revisión externa de la solicitud de excepción. Una organización de revisión externa revisa la queja que es para pedir una excepción externa. La organización de revisión externa no está afiliada ni trabaja para SFHP. La organización de revisión externa decidirá si SFHP debe cubrir el medicamento no incluido en el formulario con base en la necesidad médica. SFHP le notificará a usted y al médico que receta sobre su decisión en un plazo máximo de 72 horas. Si la solicitud original era expedita, SFHP les notificará a usted y al médico que receta en un plazo de 24 horas.

**Nota: De conformidad con la sección 1367.22 del Health and Safety Code, SFHP no limitará ni excluirá la cobertura de un medicamento si el plan de salud aprobó previamente la cobertura del medicamento para la afección médica de un afiliado y el proveedor que lo receta continúa recetándolo para la afección médica, siempre que el medicamento se recete adecuadamente y sea seguro y eficaz para tratar la afección médica del afiliado.

Otras políticas del Formulario

Política de medicamentos de marca

SFHP tiene una política genérica obligatoria y requiere la sustitución por medicamento genérico cuando está disponible un producto genérico o biológico intercambiable con calificación AB equivalente por un medicamento de marca. La dispensación de estos medicamentos de marca está permitida solo en determinados casos:

- La farmacia factura los medicamentos de marca como productos genéricos.
- La farmacia está dispensando uno (1) de los siguientes medicamentos/clases de índice terapéutico estrecho: Dilantin (fenitoína), hormonas tiroideas, Coumadin (warfarina).
- Autorización previa con documentación que indique que se probaron dos (2) medicamentos genéricos de diferentes fabricantes que no satisfacen las necesidades médicas del afiliado.

Debe enviar la solicitud de autorización previa siguiendo las instrucciones indicadas anteriormente.

Puede haber excepciones poco frecuentes a esta política. Si el medicamento de marca se incluye o se prefiere en el formulario cuando hay disponible un producto genérico o biológico intercambiable equivalente, se aplicará el costo compartido más bajo (copago para genérico).

Política de suministro diario

La política estándar de suministro diario de SFHP es un límite de suministro de recetas de 30 días para la mayoría de los medicamentos de marca y un límite de suministro de recetas de 90 días para medicamentos genéricos, con algunas excepciones. Los nuevos surtidos se permiten cuando se ha utilizado el 75% de los medicamentos de la receta médica anterior, excepto para los medicamentos opiáceos para el dolor, en cuyo caso los nuevos surtidos se permiten cuando se ha utilizado el 90% del medicamento.

Las excepciones a la política de límite de suministro de recetas para 30 días para medicamentos de marca son las siguientes:

- Se permite un suministro de hasta 12 meses para anticonceptivos y dispositivos anticonceptivos
- Se permite un suministro de hasta 100 días de tiras reactivas, lancetas, jeringas de insulina y tiras reactivas de cetonas en orina
- Se permite un suministro de hasta 90 días para determinados medicamentos utilizados en el tratamiento de afecciones crónicas. Los ejemplos de clases de medicamentos cubiertos por esta política incluyen, entre otros, medicamentos antidiabéticos, incluyendo insulina, anticonvulsivos, anticoagulantes, antidepresivos, antihiperlipidémicos, antihipertensivos, esteroides inhalados.

Las excepciones a la política de límite de suministro de 90 días para medicamentos genéricos son las siguientes:

- Se permite un suministro máximo de 30 días por surtido para todos los medicamentos opiáceos
- Se permite un suministro máximo de 14 días por surtido de medicamentos para la hepatitis C.

Política de intercambio terapéutico

Según el American College of Clinical Pharmacy (ACCP), el intercambio terapéutico se define como la dispensación de un medicamento que es terapéuticamente equivalente, pero químicamente diferente, del medicamento recetado originalmente por un médico u otro profesional autorizado. SFHP sigue la definición de intercambio terapéutico del ACCP y solo empleará el intercambio terapéutico con la aprobación del profesional que receta. Los criterios a tener en cuenta en el intercambio terapéutico incluyen la disponibilidad de fármacos dentro de una clase terapéutica, la equivalencia terapéutica, los datos de seguridad y los costos.

ENGLISH - ATTENTION: If you need help in your language, call **1(415) 547-7800** (TTY: **711**). Aids and services for people with disabilities, like documents in braille and large print, are also available. Call **1(415) 547-7800** (TTY: **711**). These services are free.

العربية (ARABIC) -

يرجى الانتباه: إذا احتجت إلى المساعدة بلغتك، فاتصل بـ **1(415) 547-7800** (TTY: **711**).
 توفر أيضاً المساعدات والخدمات للأشخاص ذوي الإعاقة، مثل المستندات المكتوبة بطريقة برييل والخط الكبير.
 اتصل بـ **1(415) 547-7800** (TTY: **711**). هذه الخدمات مجانية.

Հայերեն (ARMENIAN) - ՈՒՇԱԴՐՈՒԹՅՈՒՆ: Եթե Ձեզ օգնություն է հարկավոր Զեր լեզվով,
 զանգահարեք **1(415) 547-7800** (TTY: **711**): Կան նաև օժանդակ միջոցներ ու ծառայություններ
 հաշվանդամություն ունեցող անձանց համար, օրինակ՝ Բրայլի գրատիպով ու խոշորատառ
 տպագրված նյութեր: Զանգահարեք **1(415) 547-7800** (TTY: **711**): Այս ծառայություններն անվճար են:

ខេម (CAMBODIAN) - ចំណាំ៖ បើមួយ ត្រូវ ការជំនួយ ជាតិសាស្ត្រ របស់អ្នក សូម ទូរសព្ទទីផ្សារៗ
1(415) 547-7800 (TTY: **711**)។ ជំនួយ នឹង សេវាកម្ម ស្របតាម ជនពិភាក្សា នូចជាប័ណ្ណសាស្ត្រជាមុន្យជុំស
 ស្របតាម ជនពិភាក្សាដូច ប្រុងការសារសារជាមុន្យក្នុងជំនួយ តុលាងជាមួយ ទូរសព្ទបន្ទាល់ៗ
1(415) 547-7800 (TTY: **711**)។ សេវាកម្មទាំងនេះ គឺត្រូវការចិត្តការណ៍។

简体中文标语 (CHINESE - SIMPLIFIED) - 请注意：如果您需要以您的母语提供帮助，请致电 **1(415) 547-7800** (TTY: **711**)。另外还提供针对残疾人士的帮助和服务，例如文盲和需要较大字体阅读，也是方便可用的。请致电 **1(415) 547-7800** (TTY: **711**)。这些服务是免费的。

繁體中文 (CHINESE - TRADITIONAL) - 請注意：如果您需要以您的母語提供幫助，請致電 **1(415) 547-7800** (TTY: **711**)。另外還提供針對殘障人士的說明和服務，例如盲文和需要較大字體閱讀，也是方便可用的。請致電 **1(415) 547-7800** (TTY: **711**)。這些服務是免費的。

(FARSI - فارسی)

توجه: اگر می خواهید به زبان خود کمک دریافت کنید، با **1(415) 547-7800** (TTY: **711**) تماس بگیرید.
 کمکها و خدمات مخصوص افراد دارای معلویت، مانند نسخه های خط بربل و چاپ با حروف بزرگ، نیز
 موجود است. با **1(415) 547-7800** (TTY: **711**) تماس بگیرید. این خدمات رایگان هستند.

हिंदी (HINDI) - ध्यान दें: यदि आपको अपनी भाषा में मदद चाहिए, तो **1(415) 547-7800** (TTY: **711**)। विकलांग लोगों के लिए सहायता और सेवाएँ, जैसे ब्रेल और बड़े प्रिंट में दस्तावेज़ भी उपलब्ध हैं। **1(415) 547-7800** (TTY: **711**)। ये सेवाएँ निःशुल्क हैं।

HMOOB (HMONG) - CEEB TOOM: Yog koj xav tau kev pab txhais koj hom lus hu rau **1(415) 547-7800** (TTY: **711**). Muaj cov kev pab txhawb thiab kev pab cuam rau cov neeg xiam oob qhab, xws li puav leej muaj ua cov ntawv su thiab luam tawm ua tus ntawv loj. Hu rau **1(415) 547-7800** (TTY: **711**). Cov kev pabcuam no pub dawb.

日本語 (JAPANESE) - 注記: あなたの言語でサポートが必要な場合は、**1(415) 547-7800** (TTY: **711** までお電話ください)。また、点字や大きな活字で作成したドキュメントなど、障害をお持ちの方のための補助やサービスもご利用いただけます。**1(415) 547-7800** (TTY: **711** までお電話ください)。これらのサービスは無料です。

한국어 (KOREAN) - 주의: 자국어로 도움이 필요한 경우, **1(415) 547-7800** (TTY: 711으로 전화하십시오). 점자 및 큰 글씨로 된 문서 등 장애인을 위한 보조 도구와 서비스도 제공됩니다. **1(415) 547-7800** (TTY: 711으로 전화하십시오). 이러한 서비스는 무료입니다.

ລາວ (LAO) - ຂໍຄວນນະວັງ: ລ້າທ່ານຕົ້ນງານຄວາມຊ່ວຍເຫຼືອໃນພາສາຂອງທ່ານ, ໃຫ້ໄທໜາ
1(415) 547-7800 (TTY: 711). ການຊ່ວຍເຫຼືອ ດະກຳ: ການບໍລິການສ່າວັບຄົນຜິການເຊັ່ນ: ເອກະພານທີ່ເປັນຕົວ
ອໍານວຍອຸນນຸມ ດະກຳ ດົວຜົມຂະໜາດໃຫຍ່ ດັ່ງນັ້ນແລ້ວ. ໂທ **1(415) 547-7800** (TTY: 711). ການບໍລິການເຖິ່ງນັ້ນແລ້ວ

MIEN (MIEN) - LONGC HNYOUV JANGX LONGX OC: Beiv taux meih qiemx longc mienh tengx faan benx meih nyei waac nor douc waac daaih lorx taux **1(415) 547-7800** (TTY: 711). Liouh lorx jauv-louc tengx aengx caux nzie gong bun taux ninh mbuo wuaaic fangx mienh, beiv taux longc benx nzangc-pokc bun hluo mbiutc aengx caux aamz mborqv benx domh sou se mbenc nzoih bun longc. Douc waac daaih lorx **1(415) 547-7800** (TTY: 711). Naaiv deix gong benx wangv henh tengx oc.

ਪੰਜਾਬੀ (PUNJABI) - ਧਿਆਨ ਦਿਓ: ਜੇ ਤੁਹਾਨੂੰ ਆਪਣੀ ਭਾਸ਼ਾ ਵਿੱਚ ਮਦਦ ਦੀ ਲੋੜ ਹੈ ਤਾਂ ਕਾਲ ਕਰੋ **1(415) 547-7800** (TTY: 711 'ਤੇ ਕਾਲ ਕਰੋ)। ਅਪਾਰਸ ਲੋਕਾਂ ਲਈ
ਸਹਾਇਤਾ ਅਤੇ ਸੇਵਾਵਾਂ, ਜਿਵੇਂ ਕਿ ਬੇਲ ਅਤੇ ਮੇਟੀ ਫਾਈ ਵਿੱਚ ਦਸਤਾਵੇਜ਼, ਵੀ ਉਪਲਬਧ ਹਨ। ਕਾਲ ਕਰੋ **1(415) 547-7800** (TTY: 711 'ਤੇ ਕਾਲ ਕਰੋ)। ਇਹ ਸੇਵਾਵਾਂ ਮੁਫਤ ਹਨ।

РУССКИЙ (RUSSIAN) - ВНИМАНИЕ! Если вам нужна помощь на вашем родном языке, звоните по номеру **1(415) 547-7800** (линия TTY: 711). Также предоставляются средства и услуги для людей с ограниченными возможностями, например документы крупным шрифтом или шрифтом Брайля. Звоните по номеру **1(415) 547-7800** (линия TTY: 711). Эти услуги являются бесплатными.

ESPAÑOL (SPANISH) - ATENCIÓN: si necesita ayuda en su idioma, llame al **1(415) 547-7800** (TTY: 711). También ofrecemos asistencia y servicios para personas con discapacidades, como documentos en braille y con letras grandes. Llame al **1(415) 547-7800** (TTY: 711). Estos servicios son gratuitos.

TAGALOG (TAGALOG-FILIPINO) - ATENSIYON: Kung kailangan mo ng tulong sa iyong wika, tumawag sa **1(415) 547-7800** (TTY: 711). Mayroon ding mga tulong at serbisyo para sa mga taong may kapansanan, tulad ng mga dokumento sa braille at malaking print. Tumawag sa **1(415) 547-7800** (TTY: 711). Libre ang mga serbisyong ito.

ภาษาไทย (THAI) - โปรดทราบ: หากคุณต้องการความช่วยเหลือในภาษาของคุณ กรุณาโทรติดต่อไปที่หมายเลข **1(415) 547-7800** (TTY: 711)
นอกจากนี้ ยังพร้อมให้ความช่วยเหลือและบริการต่าง ๆ สำหรับบุคคลที่มีความพิการ เช่น เอกสารต่าง ๆ ที่เป็นอักษรเบรลล์และเอกสารที่พิมพ์ด้วยตัวอักษรขนาดใหญ่ กรุณาโทรติดต่อไปที่หมายเลข **1(415) 547-7800** (TTY: 711) บริการไม่มีค่าใช้จ่ายใด ๆ

УКРАЇНСЬКОЮ (UKRAINIAN) - УВАГА! Якщо вам потрібна допомога вашою рідною мовою, телефонуйте на номер **1(415) 547-7800** (TTY: 711). Люди з обмеженими можливостями також можуть скористатися допоміжними засобами та послугами, наприклад, отримати документи, надруковані шрифтом Брайля та великим шрифтом. Телефонуйте на номер **1(415) 547-7800** (TTY: 711). Ці послуги є безкоштовними.

TIẾNG VIỆT (VIETNAMESE) - CHÚ Ý: Nếu quý vị cần trợ giúp bằng ngôn ngữ của mình, vui lòng gọi số **1(415) 547-7800** (TTY: 711). Chúng tôi cũng hỗ trợ và cung cấp các dịch vụ dành cho người khuyết tật, như tài liệu bằng chữ nổi Braille và chữ khổ lớn (chữ hoa). Vui lòng gọi số **1(415) 547-7800** (TTY: 711). Những dịch vụ này đều là miễn phí.

AVISO DE NO DISCRIMINACIÓN

La discriminación está prohibida por ley. San Francisco Health Plan (SFHP) cumple con todas las leyes federales de derechos civiles. SFHP no discrimina, excluye ni trata a las personas de manera diferente debido a su raza, color, nacionalidad, ascendencia, religión, sexo, estado civil, género, identidad de género, orientación sexual, edad o discapacidad.

SFHP también proporciona:

- Ayuda y servicios gratuitos a las personas con discapacidades para comunicarse mejor con nosotros, como:
 - Intérpretes de lenguaje de señas calificados.
 - Información escrita en otros formatos (letra grande, audio, formatos electrónicos accesibles, otros formatos)
- Servicios de idiomas gratuitos para las personas cuyo idioma principal no es el inglés, como:
 - Intérpretes cualificados
 - Información escrita en otros idiomas.

Si necesita estos servicios, comuníquese con Servicio al Cliente de SFHP, de 8:30am a 5:30pm, de lunes a viernes, llamando al **1(415) 547-7800** o al **1(800) 288-5555** (llamada gratuita). O, si no puede escuchar o hablar bien, llame al TTY **711**.

CÓMO PRESENTAR UNA QUEJA

Si considera que SFHP no cumplió con proporcionar estos servicios o cometió discriminación de alguna otra forma, con base a la raza, color, nacionalidad, ascendencia, religión, sexo, estado civil, género, identidad de género, orientación sexual, edad o discapacidad, puede presentar una queja ante SFHP. Usted puede presentar una queja por teléfono, por escrito o por vía electrónica:

- Por teléfono: Comuníquese con SFHP de 8:30am a 5:30pm, de lunes a viernes, llamando al **1(415) 547-7800** o al **1(800) 288-5555** (llamada gratuita). O, si no puede hablar o escuchar bien, llame a la línea TTY **711** (llamada gratuita).
- Por escrito: llene un formulario de queja o escriba una carta y envíela a:
San Francisco Health Plan
P.O. Box 194247
San Francisco, CA 94119
- En persona: Visite al consultorio de su médico o el Centro de Servicio de SFHP y diga que desea presentar una queja. El Centro de Servicio de SFHP se ubica en 550 Kearny Street, Lower Level, San Francisco, CA 94108.
- Por vía electrónica: Visite el sitio web de SFHP en **sfhp.org**.

OFICINA DE DERECHOS CIVILES

Si considera que ha sido discriminado sobre la base de la raza, color, nacionalidad, edad, discapacidad o sexo, también puede presentar una queja de derechos civiles ante la U.S. Department of Health and Human Services, Office for Civil Rights, por teléfono, por escrito o por vía electrónica:

- Por teléfono: Llame al **1(800) 368-1019**. Si no puede hablar o escuchar bien, llame a la línea TTY **1(800) 537-7697**.
- Por escrito: Llene un formulario de queja o envíe una carta a:
U.S. Department of Health and Human Services
200 Independence Avenue, SW Room 509F, HHH Building
Washington, D.C. 20201

Los formularios de queja están disponibles en

<http://www.hhs.gov/ocr/office/file/index.html>.

- Por vía electrónica: Visite el portal de quejas de la Office for Civil Rights (Oficina de Derechos Civiles) en <https://ocrportal.hhs.gov/ocr/portal/lobby.jsf>.

| TIER | DESCRIPTION |
|------|--|
| 1 | Tier1 |
| 2 | Tier2 |
| 3 | Tier3 |
| TYPE | DESCRIPTION |
| QL | Quantity Limit There is a limit on the amount of this drug that is covered per prescription, or within a specific time frame. |
| CC | Clinical Criteria Your provider is required to get prior authorization before you fill your prescription, which ensures appropriate use of the selected drug. Without prior approval, we may not cover this drug. |
| ST | Step Therapy In some cases, you may be required to first try certain drugs to treat your medical condition before you move up a "step" to other drug options. |
| AL | Age Limit This prescription drug may only be covered if you meet the minimum or maximum age limit. |
| C | Custom This drug has unique restrictions. |
| HCR | Health Care Reform Products The Affordable Care Act (ACA) requires certain preventive generic products to be covered at zero dollar copay. This does not include plans that are grandfathered. |
| PA | PA Applies Your provider is required to get prior authorization before you fill your prescription, which ensures appropriate use of the selected drug. Without prior approval, we may not cover this drug. |
| QPD | Quantity Per Day Quantity Per Day. |
| \$0 | \$0 Copay \$0 Copay |

LIST OF COVERED OVER-THE-COUNTER MEDICATIONS

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| ANTI-INFECTIVES (SKIN, MUCOUS MEMBRANE) | | |
| LOCAL ANTI-INFECTIVES, MISCELLANEOUS | | |
| <i>alcohol antiseptic pads med. pad</i> | 1 | |
| ANTIANEMIA DRUGS | | |
| IRON PREPARATIONS | | |
| FEROSUL | 1 | HCR \$0 |
| FERRO-TIME | 1 | HCR \$0 |
| <i>ferrous sulfate 325(65) mg tablet</i> | 1 | HCR \$0 |
| ANTIEMETICS | | |
| ANTIHISTAMINES (GI DRUGS) | | |
| <i>meclizine hcl</i> | 1 | |
| ANTIFUNGALS (SKIN AND MUCOUS MEMBRANE) | | |
| AZOLES (SKIN AND MUCOUS MEMBRANE) | | |
| <i>clotrimazole 1 % solution</i> | 1 | QL 180 / 30 days |
| ANTIHISTAMINE DRUGS | | |
| SECOND GENERATION ANTIHISTAMINES | | |
| <i>levocetirizine dihydrochloride</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-------------------------|
| ANTIULCER AGENTS AND ACID SUPPRESSANTS | | |
| HISTAMINE H2-ANTAGONISTS | | |
| <i>cimetidine</i> | 1 | |
| <i>famotidine 20 mg tablet</i> | 1 | |
| PROTON-PUMP INHIBITORS | | |
| <i>esomeprazole magnesium 20 mg capsule dr</i> | 1 | |
| <i>lansoprazole 15 mg capsule dr</i> | 1 | |
| AUTONOMIC DRUGS | | |
| SMOKING CESSATION AGENTS | | |
| NICORETTE 2 MG LOZENGE | 1 | QPD 20.0 per day \$0 |
| <i>nicotine (7mg/24hr patch td24, 14mg/24hr patch td24, 21 mg/24hr patch td24, 21-14-7mg patch dysq)</i> | 1 | QPD 1.0 per day \$0 |
| <i>nicotine polacrilex (2 mg gum, 4 mg gum)</i> | 1 | QPD 12.0 per day \$0 |
| <i>nicotine polacrilex (2 mg lozenge, 2 mg lozng mini, 4 mg lozenge, 4 mg lozng mini)</i> | 1 | QPD 20.0 per day \$0 |
| QUIT 2 MG CHEWING GUM | 1 | QPD 12.0 per day \$0 |
| QUIT 2 MG LOZENGE | 1 | QPD 20.0 per day \$0 |
| QUIT 4 MG CHEWING GUM | 1 | QPD 12.0 per day \$0 |
| QUIT 4 MG LOZENGE | 1 | QPD 20.0 per day \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| STOP SMOKING AID | 1 | QPD \$0 | 20.0 per day |
| DEVICES | | | |
| ACCU-CHEK AVIVA SOLUTION | 1 | | |
| ACCU-CHEK FASTCLIX LANCET DRUM | 1 | | |
| ACCU-CHEK GUIDE L1-L2 CTRL SOL (NDC: 65702071310) | 1 | | |
| ACCU-CHEK GUIDE ME GLUCOSE MTR | 1 | QL | 1 / 365 days |
| ACCU-CHEK GUIDE MONITOR SYSTEM | 1 | QL | 1 / 365 days |
| ACCU-CHEK SMARTVIEW CONTRL SOL | 1 | | |
| <i>covid-19 antigen immunoassay test</i> | 2 | QL \$0 | 8 / 30 days |
| <i>covid-19 molecular nucleic acid test assay</i> | 2 | QL \$0 | 8 / 30 days |
| <i>diabetic needles</i> | 1 | | |
| <i>diabetic syringes</i> | 1 | | |
| <i>digital thermometer</i> | 1 | | |
| <i>gloves (each, package)</i> | 1 | | |
| <i>inhaler, assist devices, accessories</i> | 2 | | |
| <i>inhaler, assist device with large mask</i> | 2 | QL | 2 / 365 days |
| <i>inhaler, assist device with small mask</i> | 2 | QL | 2 / 365 days |
| <i>lancets each</i> | 1 | | |
| <i>lancing device/lancets</i> | 1 | | |
| <i>medical supply, miscellaneous each</i> | 1 | | |
| <i>nasal airflow strips strip</i> | 2 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|------------------------------------|------|-----------------------|
| <i>nebulizer</i> | 2 | QL 2 / 365 days |
| <i>nebulizer and compressor</i> | 2 | QL 2 / 365 days |
| <i>peak flow meter</i> | 2 | QL 2 / 365 days |
| <i>spirometers and accessories</i> | 2 | QL 2 / 365 days |

DIAGNOSTIC AGENTS

DIABETES MELLITUS

| | | |
|---|---|-----------------|
| ACCU-CHEK AVIVA PLUS TEST STRP (NDC: 65702040710) | 1 | QPD 4.0 per day |
| ACCU-CHEK AVIVA PLUS TEST STRP (NDC: 65702040810) | 1 | QPD 4.0 per day |
| ACCU-CHEK GUIDE TEST STRIP (NDC: 65702071110) | 1 | QPD 4.0 per day |
| ACCU-CHEK GUIDE TEST STRIP (NDC: 65702071210) | 1 | QPD 4.0 per day |
| ACCU-CHEK SMARTVIEW TEST STRIP | 2 | QPD 4.0 per day |
| <i>blood sugar diagnostic</i> | 1 | QPD 4.0 per day |

EMOLLIENTS, DEMULCENTS, AND PROTECTANTS

BASIC LOTIONS AND LINIMENTS

| | |
|-------------------------------------|---|
| <i>ammonium lactate 12 % lotion</i> | 1 |
|-------------------------------------|---|

BASIC OINTMENTS AND PROTECTANTS

| | |
|--|---|
| <i>ammonium lactate 12 % cream (g)</i> | 1 |
|--|---|

EYE, EAR, NOSE AND THROAT (EENT) PREPS.

ANTIALLERGIC AGENTS

| | |
|-------------------|---|
| ALAWAY | 1 |
| ALLERGY EYE DROPS | 1 |
| CHILDREN'S ALAWAY | 1 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| EYE ITCH RELIEF | 1 | | |
| <i>ketotifen fumarate 0.025 % drops</i> | 1 | | |
| <i>olopatadine hcl (0.1 % drops, 0.2 % drops)</i> | 3 | CC ST QPD | 0.17 per day |
| PATADAY ONCE DAILY 0.7% DROPS | 3 | CC ST | |
| WAL-ZYR 0.025% EYE DROPS | 1 | | |
| ZADITOR | 1 | | |
| HORMONES AND SYNTHETIC SUBSTITUTES | | | |
| CONTRACEPTIVES | | | |
| AFTER PILL | 1 | HCR \$0 | |
| AFTERA | 1 | \$0 | |
| ECONTRA EZ | 1 | \$0 | |
| ECONTRA ONE-STEP | 1 | \$0 | |
| HER STYLE | 1 | HCR \$0 | |
| <i>levonorgestrel</i> | 1 | \$0 | |
| MY CHOICE | 1 | \$0 | |
| MY WAY | 1 | \$0 | |
| NEW DAY | 1 | \$0 | |
| OPCICON ONE-STEP | 1 | \$0 | |
| OPILL | 2 | \$0 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| OPTION 2 | 1 | \$0 |
| TAKE ACTION | 1 | \$0 |
| INSULINS | | |
| INTERMEDIATE-ACTING INSULINS | | |
| HUMULIN N | 2 | |
| HUMULIN N KWIKPEN | 2 | |
| NOVOLIN N | 2 | |
| NOVOLIN N FLEXPEN | 2 | |
| SHORT-ACTING INSULINS | | |
| HUMULIN R | 2 | |
| NOVOLIN R | 2 | |
| NOVOLIN R FLEXPEN | 2 | |
| NONHORMONAL CONTRACEPTIVES | | |
| <i>condoms, female</i> | 1 | \$0 |
| <i>condoms, latex, lubricated</i> | 1 | \$0 |
| <i>condoms, latex, non-lubricated</i> | 1 | \$0 |
| <i>condoms, non-latex, lubricated</i> | 1 | \$0 |
| VCF (FILM, GEL) | 1 | \$0 |
| NONSTEROIDAL ANTI-INFLAMMATORY AGENTS | | |
| REVERSIBLE COX-1/COX-2 INHIBITORS | | |
| <i>ibuprofen 100 mg/5ml oral susp</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| SALICYLATES | | |
| <i>aspirin 81 mg</i> | 1 | \$0 |
| SKIN AND MUCOUS MEMBRANE AGENTS | | |
| KERATOLYTIC AGENTS | | |
| ACNE MEDICATION 5% GEL | 1 | QL 60 / 30 days |
| <i>benzoyl peroxide 5 % gel (gram)</i> | 1 | QL 60 / 30 days |
| URINE AND FECES CONTENTS | | |
| KETONES | | |
| <i>urine acetone test,strips</i> | 1 | QL 100 / 100 days |
| VITAMINS | | |
| MULTIVITAMIN PREPARATIONS | | |
| <i>prenatal with folic acid</i> | 1 | \$0 |
| VITAMIN B COMPLEX | | |
| <i>folic acid 0.4 mg tablet</i> | 1 | \$0 |
| <i>folic acid 1 mg tablet</i> | 1 | \$0 |
| MYNEPHRON | 1 | |
| NEPHRO-VITE | 1 | |
| RENA-VITE | 1 | |

LIST OF COVERED PRESCRIPTION MEDICATIONS

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| ALPHA-ADRENERGIC BLOCKING AGENT(SYMPATH) NON-SEL.ALPHA-ADRENERGIC BLOCKING AGENTS | | | |
| ERGOMAR | 2 | | |
| <i>ergotamine tartrate/caffeine</i> | 1 | | |
| SELECTIVE ALPHA-1-ADRENERGIC BLOCK.AGENT | | | |
| <i>alfuzosin hcl</i> | 1 | | |
| <i>tamsulosin hcl</i> | 1 | | |
| ANALGESICS AND ANTIPYRETICS | | | |
| OPPIOID AGONISTS (28:08) | | | |
| <i>acetaminophen with codeine 120-12mg/5 solution</i> | 1 | CC QPD | 12.0 per day |
| <i>acetaminophen with codeine phosphate (300mg-15mg tablet, 300mg-30mg tablet, 300mg-60mg tablet)</i> | 1 | CC QPD | 4.0 per day |
| <i>codeine sulfate (30 mg tablet, 60 mg tablet)</i> | 1 | CC QPD | 4.0 per day |
| <i>codeine sulfate 15 mg tablet</i> | 1 | CC | |
| ENDOCET | 1 | CC QPD | 4.0 per day |
| <i>fentanyl</i> | 3 | QL CC PA | 15 / 30 days |
| <i>hydrocodone bitartrate/acetaminophen (hydrocodone/acetaminophen 5 mg-325mg tablet, hydrocodone/acetaminophen 7.5-325 mg tablet, hydrocodone/acetaminophen 10mg-325mg tablet)</i> | 1 | CC QPD | 4.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|------------------------|
| hydrocodone/acetaminophen 2.5-325 mg tablet | 1 | QL CC QPD | 4 / day 4.0 per day |
| hydromorphone hcl (2 mg tablet, 4 mg tablet) | 1 | CC QPD | 4.0 per day |
| hydromorphone hcl 8 mg tablet | 1 | QL CC QPD | 4 / day 4.0 per day |
| morphine sulfate (10 mg/5 ml solution, 20 mg/5 ml solution, 100 mg/5ml solution) | 1 | CC QPD | 12.0 per day |
| morphine sulfate 15 mg tablet | 1 | CC QPD | 4.0 per day |
| morphine sulfate 30 mg tablet | 1 | QL CC QPD | 4 / day 4.0 per day |
| morphine sulfate (15 mg tablet er, 100 mg tablet er, 200 mg tablet er) | 1 | QL CC QPD | 3 / day 3.0 per day |
| morphine sulfate (30 mg tablet er, 60 mg tablet er) | 1 | CC QPD | 3.0 per day |
| oxycodone hcl (5 mg/5 ml solution, 20 mg/ml oral conc) | 1 | CC QPD | 12.0 per day |
| oxycodone hcl (10 mg tab er 12h, 20 mg tab er 12h, 40 mg tab er 12h, 80 mg tab er 12h) | 3 | CC PA QPD | 2.0 per day |
| oxycodone hcl (5 mg tablet, 10 mg tablet, 15 mg tablet, 20 mg tablet, 30 mg tablet) | 1 | CC QPD | 4.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|---------------|
| <i>oxycodone hcl/acetaminophen (hcl/acetaminophen 2.5-325 mg tablet, hcl/acetaminophen 5 mg-325mg tablet, hcl/acetaminophen 7.5-325 mg tablet, hcl/acetaminophen 10mg-325mg tablet)</i> | 1 | CC QPD | 4.0 per day |
| <i>oxymorphone hcl (5 mg tablet, 10 mg tablet)</i> | 1 | | |
| <i>tramadol hcl 50 mg tablet</i> | 1 | CC QPD | 8.0 per day |
| <i>tramadol hcl/acetaminophen</i> | 1 | CC QPD | 4.0 per day |
| OPIOID PARTIAL AGONISTS | | | |
| BRIXADI | 2 | | |
| <i>buprenorphine (5 mcg/hr patch tdwk, 10 mcg/hr patch tdwk)</i> | 1 | CC | |
| <i>buprenorphine hcl (2 mg tab subl, 8 mg tab subl)</i> | 1 | QL | 120 / 30 days |
| <i>buprenorphine hcl/naloxone hcl (/naloxone 2 mg-0.5mg film, /naloxone 4mg-1mg film, /naloxone 8 mg-2 mg film, /naloxone 12 mg-3 mg film)</i> | 3 | QL CC PA | 120 / 30 days |
| <i>buprenorphine hcl/naloxone hcl (/naloxone 2 mg-0.5mg tab subl, /naloxone 8 mg-2 mg tab subl)</i> | 1 | QL | 120 / 30 days |
| SUBLOCADE | 2 | | |
| ZUBSOLV | 3 | QL CC PA | 120 / 30 days |
| ANOREXIGENIC AGENTS | | | |
| AMPHETAMINE DERIVATIVES | | | |
| <i>phentermine hcl (15 mg capsule, 30 mg capsule, 37.5 mg capsule, 37.5 mg tablet)</i> | 3 | QL CC PA | 30 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|---|
| ANOREXIGENICS;RESPIRATORY,CNS STIMULANTS AMPHETAMINES | | |
| <i>dextroamphetamine sulf-saccharate/amphetamine sulf-aspartate (dextroamphetamine/amphetamine 5 mg cap er 24h, dextroamphetamine/amphetamine 5 mg tablet, dextroamphetamine/amphetamine 7.5 mg tablet, dextroamphetamine/amphetamine 10 mg cap er 24h, dextroamphetamine/amphetamine 10 mg tablet, dextroamphetamine/amphetamine 12.5 mg tablet, dextroamphetamine/amphetamine 15 mg cap er 24h, dextroamphetamine/amphetamine 15 mg tablet, dextroamphetamine/amphetamine 20 mg cap er 24h, dextroamphetamine/amphetamine 20 mg tablet, dextroamphetamine/amphetamine 25 mg cap er 24h, dextroamphetamine/amphetamine 30 mg cap er 24h, dextroamphetamine/amphetamine 30 mg tablet)</i> | 1 | QL 60 / 30 days |
| <i>dextroamphetamine sulfate (5 mg capsule er, 10 mg capsule er, 15 mg capsule er)</i> | 1 | QL 60 / 30 days |
| <i>dextroamphetamine sulfate 10 mg tablet</i> | 1 | QL 120 / 30 days AL At least 5 yrs old PA |
| <i>dextroamphetamine sulfate 5 mg tablet</i> | 1 | QL 60 / 30 days AL At least 5 yrs old PA |
| ANOREXIGENIC AGENTS | | |
| CONTRAVE | 3 | QL 120 / 30 days CC PA QPD 4.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|-----------------------------|
| RESPIRATORY AND CNS STIMULANTS | | | |
| <i>atomoxetine hcl</i> | 1 | QL | 60 / 30 days |
| <i>dexmethylphenidate hcl (2.5 mg tablet, 5 mg cpbp 50-50, 5 mg tablet, 10 mg cpbp 50-50, 10 mg tablet, 15 mg cpbp 50-50, 20 mg cpbp 50-50, 25 mg cpbp 50-50, 30 mg cpbp 50-50, 35 mg cpbp 50-50, 40 mg cpbp 50-50)</i> | 1 | QL | 60 / 30 days |
| METADATE ER | 1 | | |
| <i>methylphenidate hcl (10 mg cpbp 30-70, 10 mg cpbp 50-50, 10 mg tablet er, 18 mg tab er 24, 20 mg cpbp 30-70, 20 mg cpbp 50-50, 27 mg tab er 24, 30 mg cpbp 30-70, 30 mg cpbp 50-50, 36 mg tab er 24, 40 mg cpbp 30-70, 40 mg cpbp 50-50, 50 mg cpbp 30-70, 54 mg tab er 24, 60 mg cpbp 30-70, 60 mg cpbp 50-50)</i> | 1 | QL | 60 / 30 days |
| <i>methylphenidate hcl 10 mg/5 ml solution</i> | 1 | QL | 900 / 30 days |
| <i>methylphenidate hcl 5 mg/5 ml solution</i> | 1 | QL | 300 / 30 days |
| <i>methylphenidate hcl (5 mg tablet, 10 mg tablet, 20 mg tablet, 20 mg tablet er)</i> | 1 | QL | 90 / 30 days |
| WAKEFULNESS-PROMOTING AGENTS | | | |
| <i>armodafinil</i> | 3 | QL CC PA QPD | 90 / 90 days 1.0 per day |
| <i>modafinil (100 mg tablet, 200 mg tablet)</i> | 3 | CC PA QPD | 1.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|------------------------|
| ANTI-INFECTIVE AGENTS | | |
| ANTHELMINTICS | | |
| <i>albendazole 200 mg tablet</i> | 1 | QL 6 / 365 days |
| <i>ivermectin 3 mg tablet</i> | 1 | QL 30 / 365 days CC |
| <i>praziquantel 600 mg tablet</i> | 1 | QL 15 / 365 days |
| URINARY ANTI-INFECTIVES | | |
| <i>methenamine hippurate</i> | 1 | |
| <i>nitrofurantoin macrocrystal</i> | 1 | |
| <i>nitrofurantoin monohydrate/macrocystals</i> | 1 | |
| <i>trimethoprim 100 mg tablet</i> | 1 | |
| ANTI-INFECTIVES (EENT) | | |
| ANTI-INFECTIVES, MISCELLANEOUS (52:04) | | |
| <i>acetic acid 2 % solution</i> | 1 | |
| <i>hydrocortisone/acetic acid</i> | 1 | |
| ANTIBACTERIALS (52:04) | | |
| <i>AK-POLY-BAC</i> | 1 | |
| <i>bacitracin 500 unit/g oint. (g)</i> | 1 | |
| <i>bacitracin/polymyxin b sulfate</i> | 1 | |
| <i>ciprofloxacin hcl 0.3 % drops</i> | 1 | |
| <i>ciprofloxacin hcl/dexamethasone</i> | 1 | QL 7.5 / 30 days |
| <i>doxycycline hydiate 20 mg tablet</i> | 1 | QL 60 / 30 days |
| <i>erythromycin base 5 mg/gram oint. (g)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------------|
| <i>erythromycin base in ethanol (in 2 % gel (gram), in 2 % solution)</i> | 1 | |
| <i>gentamicin sulfate 0.3 % drops</i> | 1 | |
| <i>moxifloxacin hcl 400 mg tablet</i> | 3 | CC PA QPD 1.0 per day |
| NEO-POLYCIN | 1 | |
| NEO-POLYCIN HC | 1 | |
| <i>neomycin sulfate/bacitracin zinc/polymyxin b/hydrocortisone</i> | 1 | |
| <i>neomycin sulfate/bacitracin/polymyxin b</i> | 1 | |
| <i>neomycin sulfate/polymyxin b sulfate/gramicidin d</i> | 1 | |
| <i>neomycin sulfate/polymyxin b sulfate/hydrocortisone (neomycin/polymyxin b/hydrocort 3.5-10k-1 drops susp, neomycin/polymyxin b/hydrocort 3.5-10k-1 solution, neomycin/polymyxin b/hydrocort 3.5-10k-10 drops susp)</i> | 1 | |
| <i>neomycin/polymyxin b sulfate/dexamethasone (neomycin/polymyxin b/dexametha 0.1 % drops susp, neomycin/polymyxin b/dexametha 3.5-10k-.1 oint. (g))</i> | 1 | |
| <i>ofloxacin 0.3 % drops</i> | 1 | |
| POLYCIN | 1 | |
| <i>polymyxin b sulfate(trimethoprim</i> | 1 | |
| <i>sulfacetamide sodium (10 % drops, 10 % oint. (g))</i> | 1 | |
| <i>sulfacetamide sodium/prednisolone sodium phosphate</i> | 1 | |
| TOBRADEX EYE OINTMENT | 3 | ST |
| <i>tobramycin 0.3 % drops</i> | 1 | |
| <i>tobramycin/dexamethasone</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|--|
| ANTIFUNGALS (EENT) | | |
| NATACYN | 2 | |
| ANTIVIRALS (EENT) | | |
| <i>trifluridine 1 % drops</i> | 1 | |
| ZIRGAN | 2 | |
| ASTRINGENTS (52:04) | | |
| <i>chlorhexidine gluconate</i> | 1 | |
| PAROEX | 1 | |
| PERIOGARD | 1 | |
| ANTI-INFECTIVES (SKIN, MUCOUS MEMBRANE) | | |
| ANTIBACTERIALS (84:04) | | |
| <i>azelaic acid 15 % gel (gram)</i> | 1 | QL 50 / 30 days CC QPD 1.7 per day |
| AZELEX | 3 | CC PA QPD 1.0 per day |
| FINACEA 15% FOAM | 3 | CC PA QPD 1.7 per day |
| <i>metronidazole (0.75 % cream (g), 0.75 % gel (gram))</i> | 1 | QL 45 / 30 days |
| <i>metronidazole (1 % gel (gram), 1 % gel w/pump)</i> | 3 | QL 60 / 30 days ST |
| <i>metronidazole 0.75 % lotion</i> | 1 | QL 60 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|--|------|-----------------------|--|--|
| <i>mupirocin 2 % oint. (g)</i> | 1 | | | |
| ROSADAN (CREAM, GEL) | 1 | | | |
| <i>tetracycline hcl (250 mg capsule, 500 mg capsule)</i> | 1 | QL | 120 / 30 days | |
| ANTIVIRALS (SKIN AND MUCOUS MEMBRANE) | | | | |
| <i>acyclovir 5 % cream (g)</i> | 3 | CC PA QPD | 0.17 per day | |
| <i>acyclovir 5 % oint. (g)</i> | 3 | QL CC PA QPD | 0.5 / day 0.5 per day | |
| ASTRINGENTS, ANTI-INFECTIVE | | | | |
| <i>selenium sulfide 2.5 % lotion</i> | 1 | | | |
| <i>silver sulfadiazine 1 % cream (g)</i> | 1 | | | |
| SCABICIDES AND PEDICULICIDES | | | | |
| <i>malathion</i> | 3 | QL CC PA | 59 / fill | |
| <i>permethrin 5 % cream (g)</i> | 1 | QL CC | 120 / fill | |
| <i>spinosad</i> | 3 | C PA | Limit of 120 ml per fill equates to a limit of 1 bottle per fill | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|---|
| ANTI-INFLAMMATORY AGENTS (EENT) | | |
| CORTICOSTEROIDS (EENT) | | |
| <i>difluprednate</i> | 3 | QL 5 / 30 days ST |
| FLAC OTIC OIL | 1 | |
| <i>fluocinolone acetonide oil</i> | 1 | |
| <i>fluorometholone</i> | 1 | |
| <i>fluticasone propionate 50 mcg spray susp</i> | 1 | QL 16 / 30 days |
| PRED MILD | 3 | ST |
| <i>prednisolone acetate</i> | 1 | |
| <i>prednisolone sodium phosphate 1 % drops</i> | 1 | |
| EENT NONSTEROIDAL ANTI-INFLAM. AGENTS | | |
| <i>diclofenac sodium 0.1 % drops</i> | 1 | QL 5 / fill |
| <i>ketorolac tromethamine 0.4 % drops</i> | 1 | QL 5 / fill |
| <i>ketorolac tromethamine 0.5 % drops</i> | 1 | QL 10 / fill C Limit of 10 ml per fill |
| ANTI-INFLAMMATORY AGENTS (RESPIRATORY) | | |
| LEUKOTRIENE MODIFIERS | | |
| <i>montelukast sodium 10 mg tablet</i> | 1 | |
| MAST-CELL STABILIZERS | | |
| <i>cromolyn sodium (20 mg/2 ml ampul-neb, 20 mg/ml oral conc)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|-------------|
| ANTI-INFLAMMATORY AGENTS (SKIN, MUCOUS) | | | |
| CORTICOSTEROIDS (SKIN, MUCOUS MEMBRANE) | | | |
| ANUCORT-HC | 1 | | |
| ANUSOL-HC 25 MG SUPPOSITORY | 1 | | |
| <i>betamethasone dipropionate (0.05 % cream (g), 0.05 % lotion, 0.05 % oint. (g))</i> | 1 | QPD | 8.0 per day |
| <i>betamethasone dipropionate 0.05 % gel (gram)</i> | 1 | | |
| <i>betamethasone dipropionate/propylene glycol (betamethasone/propylene 0.05 % cream (g), betamethasone/propylene 0.05 % lotion, betamethasone/propylene 0.05 % oint. (g))</i> | 1 | QPD | 4.0 per day |
| <i>betamethasone valerate (0.1 % cream (g), 0.1 % lotion, 0.1 % oint. (g))</i> | 1 | QPD | 8.0 per day |
| <i>clobetasol propionate (0.05 % cream (g), 0.05 % gel (gram), 0.05 % oint. (g), 0.05 % shampoo, 0.05 % solution)</i> | 1 | QPD | 4.0 per day |
| CLODAN 0.05% SHAMPOO | 1 | QPD | 4.0 per day |
| <i>desoximetasone 0.25 % cream (g)</i> | 1 | QPD | 4.0 per day |
| <i>desoximetasone 0.25 % oint. (g)</i> | 1 | CC QPD | 4.0 per day |
| <i>fluocinolone acetonide 0.025 % cream (g)</i> | 1 | | |
| <i>fluocinolone acetonide 0.01 % oil</i> | 1 | QPD | 4.0 per day |
| <i>fluocinolone acetonide 0.01 % solution</i> | 1 | QPD | 6.0 per day |
| <i>fluocinolone acetonide/shower cap</i> | 1 | QPD | 4.0 per day |
| <i>fluocinonide (0.05 % cream (g), 0.05 % gel (gram), 0.05 % oint. (g), 0.05 % solution)</i> | 1 | QPD | 8.0 per day |
| <i>fluticasone propionate 0.05 % cream (g)</i> | 1 | QPD | 8.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|---------------|
| halobetasol propionate (0.05 % cream (g), 0.05 % oint. (g)) | 1 | QPD | 4.0 per day |
| HEMMOREX-HC 25 MG SUPPOSITORY | 1 | | |
| hydrocortisone 1 % cream (g) | 1 | CC | |
| hydrocortisone 2.5 % crm/pe app | 1 | QPD | 8.0 per day |
| hydrocortisone 100mg/60ml enema | 1 | | |
| hydrocortisone (2.5 % cream (g), 2.5 % lotion, 2.5 % oint. (g)) | 1 | QPD | 8.0 per day |
| hydrocortisone acetate 25 mg supp.rect | 1 | | |
| mometasone furoate (0.1 % cream (g), 0.1 % oint. (g), 0.1 % solution) | 1 | QPD | 8.0 per day |
| nystatin/triamcinolone acetonide (nystatin/triamcin 100000-0.1 cream (g), nystatin/triamcin 100000-0.1 oint. (g), nystatin/triamcinolone acet 100000-0.1 cream (g), nystatin/triamcinolone acet 100000-0.1 oint. (g)) | 1 | QL | 480 / 30 days |
| ORALONE | 1 | | |
| PROCTO-MED HC | 1 | QPD | 2.0 per day |
| PROCTOFOAM-HC | 2 | | |
| PROCTOSOL-HC | 1 | QPD | 2.0 per day |
| PROCTOZONE-HC | 1 | QPD | 2.0 per day |
| triamcinolone acetonide (0.025 % cream (g), 0.025 % lotion, 0.025 % oint. (g), 0.1 % lotion, 0.5 % cream (g), 0.5 % oint. (g)) | 1 | QPD | 8.0 per day |
| triamcinolone acetonide (0.1 % cream (g), 0.1 % oint. (g)) | 1 | QPD | 16.0 per day |
| triamcinolone acetonide 0.1 % paste (g) | 1 | | |
| TRIDERM 0.1% CREAM | 1 | QPD | 16.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|-------------|
| TRIDERM 0.5% CREAM | 1 | QPD | 8.0 per day |
| IMMUNOMODULATORY AGENTS (84:06) | | | |
| <i>tacrolimus (0.03 % oint. (g), 0.1 % oint. (g))</i> | 3 | CC ST QPD | 1.0 per day |
| JANUS KINASE INHIBITORS (84:06) | | | |
| CIBINOO | 3 | CC PA QPD | 1.0 per day |
| ANTIANEMIA DRUGS | | | |
| IRON PREPARATIONS | | | |
| NEONATAL FE | 2 | | |
| ANTIARRHYTHMIC AGENTS | | | |
| CLASS IA ANTIARRHYTHMICS | | | |
| <i>disopyramide phosphate</i> | 1 | | |
| NORPACE CR | 2 | | |
| <i>quinidine gluconate</i> | 1 | | |
| <i>quinidine sulfate (200 mg tablet, 300 mg tablet)</i> | 1 | | |
| CLASS IB ANTIARRHYTHMICS | | | |
| <i>mexiletine hcl (150 mg capsule, 200 mg capsule, 250 mg capsule)</i> | 1 | | |
| CLASS IC ANTIARRHYTHMICS | | | |
| <i>flecainide acetate</i> | 1 | | |
| <i>propafenone hcl (150 mg tablet, 225 mg tablet, 300 mg tablet)</i> | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| CLASS III ANTIARRHYTHMICS | | |
| <i>amiodarone hcl (100 mg tablet, 200 mg tablet, 400 mg tablet)</i> | 1 | |
| <i>dofetilide</i> | 1 | |
| MULTAQ | 2 | |
| PACERONE 200 MG TABLET | 1 | |
| CLASS IV ANTIARRHYTHMICS | | |
| CARTIA XT | 1 | |
| DILT-XR | 1 | |
| <i>diltiazem hcl (30 mg tablet, 60 mg cap er 12h, 60 mg tablet, 90 mg cap er 12h, 90 mg tablet, 120 mg cap er 12h, 120 mg cap er 24h, 120 mg cap er deg, 120 mg cap sa 24h, 120 mg tablet, 180 mg cap er 24h, 180 mg cap er deg, 180 mg cap sa 24h, 240 mg cap er 24h, 240 mg cap er deg, 240 mg cap sa 24h, 300 mg cap er 24h, 300 mg cap sa 24h, 360 mg cap er 24h, 360 mg cap sa 24h, 420 mg cap sa 24h)</i> | 1 | |
| TAZTIA XT (180 MG CAPSULE, 240 MG CAPSULE, 300 MG CAPSULE, 360 MG CAPSULE) | 1 | |
| TAZTIA XT 120 MG CAPSULE | 1 | QL 30 / 30 days |
| TIADYLT ER | 1 | |
| <i>verapamil hcl (40 mg tablet, 80 mg tablet, 120 mg cap24h pel, 120 mg tablet, 120 mg tablet er, 180 mg cap24h pel, 180 mg tablet er, 240 mg cap24h pel, 240 mg tablet er, 360 mg cap24h pel)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------------|
| ANTIBACTERIALS (08:12) | | |
| AMINOGLYCOSIDE ANTIBIOTICS | | |
| <i>neomycin sulfate 500 mg tablet</i> | 1 | |
| <i>tobramycin in 0.225 % sodium chloride</i> | 3 | CC PA QPD 5.0 per day |
| QUINOLONE ANTIBIOTICS | | |
| <i>ciprofloxacin hcl (100 mg tablet, 250 mg tablet, 500 mg tablet, 750 mg tablet)</i> | 1 | |
| <i>levofloxacin (250 mg tablet, 500 mg tablet, 750 mg tablet)</i> | 1 | |
| SULFONAMIDE ANTIBIOTICS (SYSTEMIC) | | |
| <i>sulfadiazine 500 mg tablet</i> | 1 | |
| <i>sulfamethoxazole/trimethoprim (sulfamethoxazole/trimethoprim 400mg-80mg tablet, sulfamethoxazole/trimethoprim 800-160 mg tablet)</i> | 1 | |
| <i>sulfasalazine (500 mg tablet, 500 mg tablet dr)</i> | 1 | |
| TETRACYCLINE ANTIBIOTICS | | |
| AVIDOXY | 1 | |
| <i>doxycycline hydiate (50 mg capsule, 100 mg capsule, 100 mg tablet)</i> | 1 | QL 60 / 30 days |
| <i>doxycycline monohydrate (50 mg capsule, 100 mg capsule, 100 mg tablet)</i> | 1 | QL 60 / 30 days |
| LYMEPAK | 1 | QL 60 / 30 days |
| <i>minocycline hcl (50 mg capsule, 75 mg capsule, 100 mg capsule)</i> | 1 | QL 60 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|---|
| MONDOXYNE NL 100 MG CAPSULE | 1 | |
| ANTIBACTERIALS, MISCELLANEOUS | | |
| GLYCOPEPTIDE ANTIBIOTICS | | |
| <i>vancomycin hcl (125 mg capsule, 250 mg capsule)</i> | 1 | |
| LINCOMYCIN ANTIBIOTICS | | |
| CLEOCIN 100 MG VAGINAL OVULE | 2 | |
| CLINDACIN P | 1 | |
| <i>clindamycin hcl (75 mg capsule, 150 mg capsule, 300 mg capsule)</i> | 1 | |
| <i>clindamycin phosphate (1 % gel (gram), 1 % lotion, 1 % med. swab, 1 % solution, 2 % cream/applicator)</i> | 1 | |
| OXAZOLIDINONE ANTIBIOTICS | | |
| <i>linezolid 600 mg tablet</i> | 1 | |
| RIFAMYCIN ANTIBIOTICS | | |
| XIFAXAN | 3 | CC ST QPD 3.0 per day |
| ANTICHOLINERGIC AGENTS | | |
| ANTIMUSCARINICS/ANTISPASMODICS | | |
| ATROVENT HFA | 2 | QPD 0.9 per day |
| BEVESPI AEROSPHERE | 2 | QPD 0.36 per day |
| <i>chlordiazepoxide/clidinium bromide</i> | 1 | |
| COMBIVENT RESPIMAT | 2 | QPD 0.2 per day |
| <i>dicyclomine hcl (10 mg capsule, 10 mg/5 ml solution, 20 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| <i>glycopyrrolate (1 mg tablet, 2 mg tablet)</i> | 1 | | |
| <i>hyoscyamine sulfate (0.125 mg tab subl, 0.125 mg tablet, 0.375 mg tab er 12h)</i> | 1 | | |
| INCRUSE ELLIPTA | 2 | QPD | 1.0 per day |
| <i>ipratropium bromide 0.2 mg/ml solution</i> | 1 | QPD | 11.0 per day |
| <i>ipratropium bromide/albuterol sulfate</i> | 1 | QPD | 19.0 per day |
| OSCIMIN | 1 | | |
| OSCIMIN SL | 1 | | |
| <i>scopolamine</i> | 1 | QL | 4 per fill |
| SPIRIVA RESPIMAT | 2 | QPD | 0.14 per day |
| STIOLTO RESPIMAT | 2 | QPD | 0.14 per day |
| SYMAX-SL | 1 | | |
| SYMAX-SR | 1 | | |
| TRELEGY ELLIPTA | 3 | ST QPD | 2.0 per day |
| ANTICOAGULANTS | | | |
| COUMARIN DERIVATIVES | | | |
| JANTOVEN | 1 | | |
| <i>warfarin sodium (1 mg tablet, 2 mg tablet, 2.5 mg tablet, 3 mg tablet, 4 mg tablet, 5 mg tablet, 6 mg tablet, 7.5 mg tablet, 10 mg tablet)</i> | 1 | | |
| DIRECT FACTOR XA INHIBITORS | | | |
| ELIQUIS DVT-PE TREAT START 5MG | 2 | QL | 74 / 30 days |
| ELIQUIS (2.5 MG TABLET, 5 MG TABLET) | 2 | QL | 60 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|-----------------------------|
| rivaroxaban 1 mg/ml susp recon | 1 | QPD | 30.0 per day |
| rivaroxaban 2.5 mg tablet | 1 | QPD | 2.0 per day |
| XARELTO 1 MG/ML SUSPENSION | 2 | QPD | 30.0 per day |
| XARELTO DVT-PE TREAT START 30D | 2 | QL | 51 / 30 days |
| XARELTO (2.5 MG TABLET, 15 MG TABLET, 20 MG TABLET) | 2 | QL | 60 / 30 days |
| XARELTO 10 MG TABLET | 2 | QL | 30 / 30 days |
| DIRECT THROMBIN INHIBITORS | | | |
| dabigatran etexilate mesylate (75 mg capsule, 150 mg capsule) | 3 | QL CC PA QPD | 60 / 30 days 2.0 per day |
| dabigatran etexilate mesylate 110 mg capsule | 3 | CC PA QPD | 2.0 per day |
| HEPARINS | | | |
| enoxaparin sodium (100 mg/ml syringe, 150 mg/ml syringe) | 1 | QL | 60 / 30 days |
| enoxaparin sodium (30mg/0.3ml syringe, 300 mg/3ml vial, 300mg/3ml vial) | 1 | QL | 18 / 30 days |
| enoxaparin sodium (80mg/0.8ml syringe, 120mg/.8ml syringe) | 1 | QL | 48 / 30 days |
| enoxaparin sodium 40mg/0.4ml syringe | 1 | QL | 24 / 30 days |
| enoxaparin sodium 60mg/0.6ml syringe | 1 | QL | 36 / 30 days |
| heparin sodium,porcine (10000/ml vial, 20000/ml vial) | 1 | | |
| heparin sodium,porcine in 0.45 % sodium chloride (in 12500/250 iv soln, in 25000/250 iv soln) | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| ANTICONVULSANTS | | |
| ANTICONVULSANTS, MISCELLANEOUS | | |
| <i>carbamazepine (100 mg cpmp 12hr, 100 mg tab chew, 100 mg tab er 12h, 200 mg cpmp 12hr, 200 mg tab er 12h, 200 mg tablet, 300 mg cpmp 12hr, 400 mg tab er 12h)</i> | 1 | |
| EPITOL | 1 | |
| <i>lamotrigine (25 mg tablet, 100 mg tablet, 150 mg tablet, 200 mg tablet)</i> | 1 | |
| <i>levetiracetam (250 mg tablet, 500 mg tablet, 750 mg tablet, 1000 mg tablet)</i> | 1 | |
| ROWEEPRA | 1 | |
| SUBVENITE | 1 | |
| <i>topiramate (25 mg tablet, 50 mg tablet, 100 mg tablet, 200 mg tablet)</i> | 1 | |
| BARBITURATES (ANTICONVULSANTS) | | |
| <i>primidone (50 mg tablet, 250 mg tablet)</i> | 1 | |
| BENZODIAZEPINES (ANTICONVULSANTS) | | |
| <i>clobazam (10 mg tablet, 20 mg tablet)</i> | 1 | |
| <i>clonazepam (0.5 mg tablet, 1 mg tablet, 2 mg tablet)</i> | 1 | QL 60 / 30 days |
| GABA-MEDIATED ANTICONVULSANTS | | |
| <i>divalproex sodium (125 mg tablet dr, 250 mg tab er 24h, 250 mg tablet dr, 500 mg tab er 24h, 500 mg tablet dr)</i> | 1 | |
| <i>gabapentin (100 mg capsule, 300 mg capsule, 400 mg capsule, 600 mg tablet, 800 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|----------------|
| <i>pregabalin (25 mg capsule, 50 mg capsule, 75 mg capsule, 100 mg capsule, 150 mg capsule, 200 mg capsule, 225 mg capsule, 300 mg capsule)</i> | 1 | QL | 2 / day |
| <i>valproic acid 250 mg capsule</i> | 1 | | |
| HYDANTOINS | | | |
| DILANTIN 100 MG CAPSULE | 2 | | |
| PHENYTEK | 2 | | |
| <i>phenytoin sodium extended</i> | 1 | | |
| ION CHANNEL INHIBITION AGENTS | | | |
| <i>lacosamide (50 mg tablet, 100 mg tablet, 150 mg tablet, 200 mg tablet)</i> | 1 | | |
| <i>oxcarbazepine (150 mg tablet, 300 mg tablet, 600 mg tablet)</i> | 1 | | |
| <i>zonisamide (25 mg capsule, 50 mg capsule, 100 mg capsule)</i> | 1 | | |
| SUCCINIMIDES | | | |
| <i>ethosuximide 250 mg capsule</i> | 1 | | |
| ANTIDEPRESSANTS | | | |
| ANTIDEPRESSANTS, MISCELLANEOUS | | | |
| <i>bupropion hcl 150 mg tab er 12h</i> | 1 | \$0 | |
| <i>bupropion hcl (75 mg tablet, 100 mg tab sr 12h, 100 mg tablet, 150 mg tab er 24h, 150 mg tab sr 12h, 200 mg tab sr 12h, 300 mg tab er 24h)</i> | 1 | | |
| ZURZUVAE | 3 | QL CC PA | 28 per 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| SEL.SEROTONIN,NOREPI REUPTAKE INHIBITOR | | |
| <i>desvenlafaxine succinate</i> | 1 | |
| <i>duloxetine hcl (20 mg capsule dr, 30 mg capsule dr, 40 mg capsule dr, 60 mg capsule dr)</i> | 1 | |
| <i>venlafaxine hcl (25 mg tablet, 37.5 mg cap er 24h, 37.5 mg tablet, 50 mg tablet, 75 mg cap er 24h, 75 mg tablet, 100 mg tablet, 150 mg cap er 24h)</i> | 1 | |
| SELECTIVE-SEROTONIN REUPTAKE INHIBITORS | | |
| <i>citalopram hydrobromide (10 mg tablet, 20 mg tablet, 40 mg tablet)</i> | 1 | |
| <i>escitalopram oxalate (5 mg tablet, 10 mg tablet, 20 mg tablet)</i> | 1 | |
| <i>fluoxetine hcl (10 mg capsule, 10 mg tablet, 20 mg capsule, 20 mg tablet, 40 mg capsule, 60 mg tablet)</i> | 1 | |
| <i>fluvoxamine maleate (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| <i>paroxetine hcl (10 mg tablet, 20 mg tablet, 30 mg tablet, 40 mg tablet)</i> | 1 | |
| <i>sertraline hcl (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| SEROTONIN MODULATORS | | |
| <i>mirtazapine (7.5 mg tablet, 15 mg tab rapdis, 15 mg tablet, 30 mg tab rapdis, 30 mg tablet, 45 mg tab rapdis, 45 mg tablet)</i> | 1 | |
| <i>nefazodone hcl</i> | 1 | |
| <i>trazodone hcl (50 mg tablet, 100 mg tablet, 150 mg tablet, 300 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------------|
| TRICYCLICS, OTHER NOREPI-RU INHIBITORS | | |
| <i>amitriptyline hcl (10 mg tablet, 25 mg tablet, 50 mg tablet, 75 mg tablet, 100 mg tablet, 150 mg tablet)</i> | 1 | |
| <i>clomipramine hcl (25 mg capsule, 50 mg capsule, 75 mg capsule)</i> | 1 | |
| <i>desipramine hcl (10 mg tablet, 25 mg tablet, 50 mg tablet, 75 mg tablet, 100 mg tablet, 150 mg tablet)</i> | 1 | |
| <i>doxepin hcl (10 mg capsule, 25 mg capsule, 50 mg capsule, 75 mg capsule, 100 mg capsule, 150 mg capsule)</i> | 1 | |
| <i>imipramine hcl (10 mg tablet, 25 mg tablet, 50 mg tablet)</i> | 1 | |
| <i>nortriptyline hcl (10 mg capsule, 25 mg capsule, 50 mg capsule, 75 mg capsule)</i> | 1 | |
| ANTIDIABETIC AGENTS | | |
| ALPHA-GLUCOSIDASE INHIBITORS | | |
| <i>acarbose (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| BIGUANIDES | | |
| <i>metformin hcl (500 mg tab er 24h, 500 mg tablet, 750 mg tab er 24h, 850 mg tablet, 1000 mg tablet)</i> | 1 | |
| DIPEPTIDYL PEPTIDASE-4(DPP-4) INHIBITORS | | |
| <i>alogliptin benzoate</i> | 3 | CC ST QPD 1.0 per day |
| <i>alogliptin benzoate/metformin hcl</i> | 3 | CC ST QPD 2.0 per day |
| <i>alogliptin benzoate/pioglitazone hcl</i> | 3 | CC ST |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|--|------|-----------------------|----|-------------------|
| INCRETIN MIMETICS | | | | |
| MOUNJARO | 3 | CC | ST | QPD 0.08 per day |
| OZEMPIC | 3 | CC | ST | QPD 0.108 per day |
| RYBELSUS | 3 | CC | ST | QPD 1.0 per day |
| SAXENDA | 3 | CC | PA | QPD 0.5 per day |
| WEGOVY (0.25 MG/0.5 ML PEN, 0.5 MG/0.5 ML PEN, 1 MG/0.5 ML PEN) | 3 | CC | PA | QPD 0.072 per day |
| WEGOVY (1.7 MG/0.75 ML PEN, 2.4 MG/0.75 ML PEN) | 3 | CC | PA | QPD 0.108 per day |
| ZEPBOUND (2.5 MG/0.5 ML PEN, 5 MG/0.5 ML PEN, 7.5 MG/0.5 ML PEN, 10 MG/0.5 ML PEN, 12.5 MG/0.5 ML PEN, 15 MG/0.5 ML PEN) | 3 | CC | PA | QPD 0.8 per day |
| MEGLITINIDES | | | | |
| <i>nateglinide</i> | 1 | | | |
| <i>repaglinide</i> | 1 | | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| SODIUM-GLUC COTRANSPORT 2 (SGLT2) INHIB | | |
| FARXIGA | 2 | QPD 1.0 per day |
| GLYXAMBI | 2 | QPD 1.0 per day |
| JARDIANCE | 2 | QPD 1.0 per day |
| SYNJARDY | 2 | QPD 2.0 per day |
| SYNJARDY XR (5-1,000 MG TABLET, 10-1,000 MG TABLET, 12.5-1,000 MG TAB) | 2 | QPD 2.0 per day |
| SYNJARDY XR 25-1,000 MG TABLET | 2 | QPD 1.0 per day |
| TRIJARDY XR | 2 | QPD 1.0 per day |
| XIGDUO XR (10 MG-1,000 MG TAB, 10 MG-500 MG TABLET) | 2 | QPD 1.0 per day |
| XIGDUO XR (2.5 MG-1,000 MG TAB, 5 MG-1,000 MG TABLET, 5 MG-500 MG TABLET) | 2 | QPD 2.0 per day |
| SULFONYLUREAS | | |
| <i>glimepiride (1 mg tablet, 2 mg tablet, 4 mg tablet)</i> | 1 | |
| <i>glipizide (2.5 mg tab er 24, 5 mg tab er 24, 5 mg tablet, 10 mg tab er 24, 10 mg tablet)</i> | 1 | |
| <i>glipizide/metformin hcl</i> | 1 | |
| <i>glyburide (1.25 mg tablet, 2.5 mg tablet, 5 mg tablet)</i> | 1 | |
| <i>glyburide/metformin hcl</i> | 1 | |
| THIAZOLIDINEDIONES | | |
| <i>pioglitazone hcl</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| ANTIDOTE THERAPEUTICS | | |
| ALCOHOL DETERRENTS (91:02) | | |
| <i>acamprosate calcium</i> | 1 | QL 120 / 30 days |
| <i>disulfiram (250 mg tablet, 500 mg tablet)</i> | 1 | |
| ANTIDOTES (91:04) | | |
| ACETAMINOPHEN ANTIDOTE | | |
| <i>acetylcysteine (100 mg/ml vial, 200 mg/ml vial)</i> | 1 | |
| CHEMOTHERAPY ANTIDOTES/PROTECTANTS | | |
| ELMIRON | 2 | |
| <i>leucovorin calcium (5 mg tablet, 10 mg tablet, 15 mg tablet, 25 mg tablet)</i> | 1 | |
| ANTIEMETICS | | |
| 5-HT3 RECEPTOR ANTAGONISTS | | |
| <i>granisetron hcl 1 mg tablet</i> | 1 | CC QPD 2.0 per day |
| <i>ondansetron 4 mg tab rapdis</i> | 1 | QL 180 / 30 days |
| <i>ondansetron 8 mg tab rapdis</i> | 1 | QL 90 / 30 days |
| <i>ondansetron hcl 4 mg tablet</i> | 1 | QL 180 / 30 days |
| <i>ondansetron hcl 8 mg tablet</i> | 1 | QL 90 / 30 days |
| ANTIHISTAMINES (GI DRUGS) | | |
| COMPRO | 1 | |
| <i>meclizine hcl (12.5 mg tablet, 25 mg tablet)</i> | 1 | |
| <i>prochlorperazine</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|-------------|
| <i>prochlorperazine maleate (5 mg tablet, 10 mg tablet)</i> | 1 | | |
| NEUROKININ-1 RECEPTOR ANTAGONISTS | | | |
| AKYNZEO 300-0.5 MG CAPSULE | 3 | QL CC PA | 2 / 30 days |
| <i>aprepitant (80 mg capsule, 125 mg capsule, 125mg-80mg cap ds pk)</i> | 3 | QL CC PA | 6 / 30 days |
| <i>aprepitant 40 mg capsule</i> | 3 | QL CC PA | 1 per fill |
| ANTIFUNGAL (SYSTEMIC) | | | |
| ANTIFUNGALS, MISCELLANEOUS | | | |
| <i>griseofulvin ultramicrosize (125 mg tablet, 250 mg tablet)</i> | 1 | | |
| <i>griseofulvin, microsize (125 mg/5ml oral susp, 500 mg tablet)</i> | 1 | | |
| AZOLE ANTIFUNGALS | | | |
| <i>fluconazole (10 mg/ml susp recon, 40 mg/ml susp recon, 50 mg tablet, 100 mg tablet, 150 mg tablet, 200 mg tablet)</i> | 1 | | |
| <i>itraconazole 100 mg capsule</i> | 1 | | |
| <i>voriconazole 200 mg/5ml susp recon</i> | 3 | CC PA | |
| <i>voriconazole 200 mg tablet</i> | 3 | CC PA | |
| | | QPD | 2.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|----------------|
| <i>voriconazole 50 mg tablet</i> | 3 | CC PA QPD | 4.0 per day |
| ANTIFUNGALS (SKIN AND MUCOUS MEMBRANE) | | | |
| ALLYLAMINES (SKIN AND MUCOUS MEMBRANE) | | | |
| <i>terbinafine hcl 250 mg tablet</i> | 1 | QL | 180 / 365 days |
| AZOLES (SKIN AND MUCOUS MEMBRANE) | | | |
| <i>clotrimazole 1 % solution</i> | 1 | QL | 180 / 30 days |
| <i>clotrimazole 10 mg troche</i> | 1 | | |
| <i>clotrimazole/betamethasone dip 1 %-0.05 % cream (g)</i> | 1 | QL | 180 / 30 days |
| <i>econazole nitrate 1 % cream (g)</i> | 1 | QL | 340 / 30 days |
| GYNIAZOLE 1 | 1 | | |
| <i>ketoconazole (2 % cream (g), 2 % shampoo, 200 mg tablet)</i> | 1 | | |
| <i>miconazole nitrate 200 mg supp.vag</i> | 1 | | |
| <i>terconazole (0.4 % cream/appl, 0.8 % cream/appl, 80 mg supp.vag)</i> | 1 | | |
| HYDROXYPYRIDONES (SKIN, MUCOUS MEMBRANE) | | | |
| CICLODAN (0.77% CREAM, 8% SOLUTION) | 1 | | |
| <i>ciclopirox 1 % shampoo</i> | 1 | QL | 120 / 30 days |
| <i>ciclopirox 8 % solution</i> | 1 | QL | 0.22 / day |
| <i>ciclopirox olamine 0.77 % cream (g)</i> | 1 | QL | 90 / 30 days |
| POLYENES (SKIN AND MUCOUS MEMBRANE) | | | |
| KLAYESTA | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|----------------|
| NYAMYC | 1 | | |
| <i>nystatin (100000/g cream (g), 100000/g oint. (g))</i> | 1 | QL | 120 / 30 days |
| <i>nystatin 100000/g powder</i> | 1 | QL | 1280 / 30 days |
| <i>nystatin (500k unit tablet, 100000/ml oral susp)</i> | 1 | | |
| ANTIGLAUCOMA AGENTS | | | |
| ALPHA-ADRENERGIC AGONISTS (EENT) | | | |
| <i>brimonidine tartrate (0.1 % drops, 0.15 % drops, 0.2 % drops)</i> | 1 | | |
| <i>brimonidine tartrate/timolol maleate</i> | 1 | | |
| BETA-ADRENERGIC BLOCKING AGENTS (EENT) | | | |
| <i>betaxolol hcl 0.5 % drops</i> | 1 | | |
| <i>dorzolamide hcl/timolol maleate</i> | 1 | | |
| <i>levobunolol hcl</i> | 1 | | |
| <i>timolol maleate (0.25 % drops, 0.5 % drops)</i> | 1 | | |
| CARBONIC ANHYDRASE INHIBITORS (EENT) | | | |
| <i>acetazolamide (125 mg tablet, 250 mg tablet, 500 mg capsule er)</i> | 1 | | |
| <i>dorzolamide hcl</i> | 1 | | |
| MIOTICS | | | |
| <i>pilocarpine hcl (1 % drops, 2 % drops, 4 % drops)</i> | 1 | | |
| PROSTAGLANDIN ANALOGS | | | |
| <i>bimatoprost 0.03 % drops</i> | 1 | | |
| <i>latanoprost 0.005 % drops</i> | 1 | | |
| <i>travoprost</i> | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| ANTIHEMORRHAGIC AGENTS | | |
| HEMOSTATICS | | |
| ALPHANATE (1,000-400 UNIT VIAL, 1,500-600 UNIT VIAL) | 2 | |
| ALPHANINE SD 500 UNIT VIAL | 2 | |
| HUMATE-P 1,200 UNIT VWF:RCO | 2 | |
| <i>tranexamic acid 650 mg tablet</i> | 1 | QL 30 / 30 days |
| XYNTHA 1,000 UNIT KIT | 2 | |
| XYNTHA SOLOFUSE (UNIT KIT, UNIT SYR) | 2 | |
| ANTIHISTAMINE DRUGS | | |
| SECOND GENERATION ANTIHISTAMINES | | |
| <i>desloratadine 5 mg tablet</i> | 1 | |
| <i>levocetirizine dihydrochloride 5 mg tablet</i> | 1 | |
| ANTIHYPOLYCEMIC AGENTS | | |
| GLYCOGENOLYTIC AGENTS | | |
| BAQSIMI | 2 | |
| GLUCAGON EMERGENCY KIT | 1 | |
| ANTILIPIDEMIC AGENTS | | |
| ANTILIPIDEMIC AGENTS, MISCELLANEOUS | | |
| <i>niacin 500 mg tablet</i> | 1 | |
| NIACOR | 1 | |
| BILE ACID SEQUESTRANTS | | |
| <i>cholestyramine (4 g powd pack, 4 g powder, powder)</i> | 1 | |
| <i>cholestyramine (with sugar) (4 g powd pack, 4 g powder)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|------------------------------|
| <i>colestipol hcl 1 g tablet</i> | 1 | |
| PREVALITE (PACKET, POWDER) | 1 | |
| CHOLESTEROL ABSORPTION INHIBITORS | | |
| <i>ezetimibe</i> | 1 | |
| FIBRIC ACID DERIVATIVES | | |
| <i>fenofibrate (54 mg tablet, 160 mg tablet)</i> | 1 | |
| <i>fenofibrate nanocrystallized</i> | 1 | |
| <i>fenofibrate, micronized (43 mg capsule, 67 mg capsule, 130 mg capsule, 134 mg capsule, 200 mg capsule)</i> | 1 | |
| <i>fenofibric acid (choline)</i> | 1 | |
| <i>gemfibrozil 600 mg tablet</i> | 1 | |
| HMG-COA REDUCTASE INHIBITORS | | |
| <i>atorvastatin calcium</i> | 1 | \$0 |
| <i>lovastatin (10 mg tablet, 20 mg tablet, 40 mg tablet)</i> | 1 | \$0 |
| <i>pravastatin sodium</i> | 1 | \$0 |
| <i>rosuvastatin calcium</i> | 1 | \$0 |
| <i>simvastatin (5 mg tablet, 10 mg tablet, 20 mg tablet, 40 mg tablet, 80 mg tablet)</i> | 1 | \$0 |
| PCSK9 INHIBITORS | | |
| PRALUENT PEN | 3 | CC PA QPD 0.07 per day |
| REPATHA PUSHTRONEX | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|-------------------------|
| REPATHA SURECLICK | 3 | CC | PA |
| REPATHA SYRINGE | 3 | CC | PA |
| ANTIMETABOLITES, IMMUNOSUPPRESS THERAPY | | | |
| ANTIMETABOLITES, IMMUNOSUPP THERAPY MISC | | | |
| <i>azathioprine (50 mg tablet, 75 mg tablet, 100 mg tablet)</i> | 1 | | |
| <i>mycophenolate mofetil (250 mg capsule, 500 mg tablet)</i> | 1 | | |
| <i>mycophenolate sodium</i> | 1 | | |
| ANTIMIGRAINE AGENTS | | | |
| CALCITONIN GENE-RELATED PEPTIDE ANTAG. | | | |
| AIMOVIG AUTOINJECTOR | 3 | CC | PA QPD 0.07 per day |
| EMGALITY PEN | 3 | CC | PA QPD 0.036 per day |
| EMGALITY 120 MG/ML SYRINGE | 3 | CC | PA QPD 0.036 per day |
| EMGALITY SYRINGE (100 MG/ML SYR(1 OF 3), 300 MG (100 MG X3SYR)) | 3 | QL CC PA | max 9/180 days |
| QULIPTA | 3 | CC PA QPD | 1.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|-----------------------------|
| SELECTIVE SEROTONIN AGONISTS | | | |
| <i>naratriptan hcl</i> | 3 | QL CC ST QPD | 36 / 30 days 1.2 per day |
| <i>rizatriptan benzoate (5 mg tab rapdis, 5 mg tablet, 10 mg tab rapdis, 10 mg tablet)</i> | 1 | QPD | 1.2 per day |
| <i>sumatriptan (5 mg spray, 20 mg spray)</i> | 3 | QL CC PA | 6 / 30 days |
| <i>sumatriptan succinate (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | QL QPD | 36 / 30 days 1.2 per day |
| ANTIMYCOBACTERIALS | | | |
| ANTILEPROSY AGENTS | | | |
| <i>dapsone (25 mg tablet, 100 mg tablet)</i> | 1 | | |
| ANTITUBERCULOSIS AGENTS | | | |
| <i>cycloserine 250 mg capsule</i> | 1 | | |
| <i>ethambutol hcl</i> | 1 | | |
| <i>isoniazid (100 mg tablet, 300 mg tablet)</i> | 1 | | |
| PASER | 2 | | |
| <i>pretomanid</i> | 3 | CC PA QPD | 1.0 per day |
| PRIFTIN | 2 | | |
| <i>pyrazinamide 500 mg tablet</i> | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|----|-----------------|
| rifabutin | 1 | | | |
| rifampin (150 mg capsule, 300 mg capsule) | 1 | | | |
| SIRTURO 100 MG TABLET | 3 | CC | PA | QPD 0.9 per day |
| SIRTURO 20 MG TABLET | 3 | CC | PA | QPD 4.3 per day |
| TRECATOR | 2 | | | |
| ANTINEOPLASTIC AGENTS | | | | |
| abiraterone acetate 250 mg tablet | 3 | CC | PA | |
| ABIRTEGA | 3 | CC | PA | |
| AKEEGA | 3 | CC | PA | |
| ALECENSA | 3 | CC | PA | |
| ALUNBRIG (30 MG TABLET, 90 MG TABLET, 90 MG-180 MG TAB PACK, 180 MG TABLET) | 3 | CC | PA | |
| anastrozole 1 mg tablet | 1 | \$0 | | |
| AUGTYRO | 3 | CC | PA | |
| AVMAPKI | 3 | CC | PA | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| AVMAPKI-FAKZYNJA | 3 | CC PA |
| AYVAKIT | 3 | CC PA |
| BALVERSA | 3 | CC PA |
| BESREMI | 3 | CC PA |
| <i>bexarotene 75 mg capsule</i> | 3 | CC PA |
| <i>bicalutamide</i> | 1 | |
| BOSULIF (100 MG TABLET, 400 MG TABLET, 500 MG TABLET) | 3 | CC PA |
| BRAFTOVI | 3 | CC PA |
| BRUKINSA | 3 | CC PA |
| CABOMETYX | 3 | CC PA |
| CALQUENCE | 3 | CC PA |
| <i>capecitabine</i> | 3 | CC PA |
| CAPRELSA | 3 | CC PA |
| COMETRIQ | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| COPIKTRA | 3 | CC PA |
| COTELLIC | 3 | CC PA |
| <i>cyclophosphamide (25 mg capsule, 50 mg capsule)</i> | 3 | CC PA |
| DANZITEN | 3 | CC PA |
| <i>dasatinib</i> | 3 | CC PA |
| DAURISMO | 3 | CC PA |
| EMCYT | 3 | CC PA |
| ENSACOVE | 3 | CC PA |
| ERIVEDGE | 3 | CC PA |
| ERLEADA | 3 | CC PA |
| <i>erlotinib hcl (25 mg tablet, 100 mg tablet, 150 mg tablet)</i> | 3 | CC PA |
| <i>everolimus (0.25 mg tablet, 0.5 mg tablet, 0.75 mg tablet, 1 mg tablet)</i> | 1 | |
| <i>everolimus (2 mg tab susp, 2.5 mg tablet, 3 mg tab susp, 5 mg tab susp, 5 mg tablet, 7.5 mg tablet, 10 mg tablet)</i> | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| <i>exemestane</i> | 1 | \$0 |
| EXKIVITY | 3 | CC PA |
| FAKZYNJA | 3 | CC PA |
| FARYDAK | 3 | CC PA |
| FOTIVDA | 3 | CC PA |
| FRUZAQLA | 3 | CC PA |
| <i>gefitinib</i> | 3 | CC PA |
| GILOTRIF | 3 | CC PA |
| GLEOSTINE | 3 | CC PA |
| GOMEKLI (1 MG CAPSULE, 1 MG TABLET FOR SUSP, 2 MG CAPSULE) | 3 | CC PA |
| HYCAMTIN (0.25 MG CAPSULE, 1 MG CAPSULE) | 3 | CC PA |
| <i>hydroxyurea 500 mg capsule</i> | 1 | |
| IBRANCE (75 MG CAPSULE, 75 MG TABLET, 100 MG CAPSULE, 100 MG TABLET, 125 MG CAPSULE, 125 MG TABLET) | 3 | CC PA |
| IBTROZI | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| ICLUSIG | 3 | CC PA |
| IDHIFA | 3 | CC PA |
| <i>imatinib mesylate (100 mg tablet, 400 mg tablet)</i> | 3 | CC PA |
| IMBRUVICA (70 MG CAPSULE, 140 MG CAPSULE, 140 MG TABLET, 280 MG TABLET, 420 MG TABLET) | 3 | CC PA |
| INLYTA | 3 | CC PA |
| INQOVI | 3 | CC PA |
| INREBIC | 3 | CC PA |
| ITOVEBI | 3 | CC PA |
| IWILFIN | 3 | CC PA |
| JAKAFI | 3 | CC PA |
| JAYPIRCA | 3 | CC PA |
| KISQALI | 3 | CC PA |
| KOSELUGO | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--------------------------------|------|-----------------------|
| KRAZATI | 3 | CC PA |
| <i>lapatinib ditosylate</i> | 3 | CC PA |
| LAZCLUZE | 3 | CC PA |
| <i>lenalidomide</i> | 3 | CC PA |
| LENVIMA | 3 | CC PA |
| <i>letrozole 2.5 mg tablet</i> | 1 | \$0 |
| LEUKERAN | 3 | CC PA |
| LONSURF | 3 | PA |
| LORBRENA | 3 | CC PA |
| LUMAKRAS | 3 | CC PA |
| LYNPARZA | 3 | CC PA |
| LYSODREN | 3 | CC PA |
| LYTGOBI | 3 | CC PA |
| MATULANE | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| MEKINIST (0.5 MG TABLET, 2 MG TABLET) | 3 | CC | PA |
| MEKTOVI | 3 | CC | PA |
| <i>melphalan</i> | 1 | | |
| <i>mercaptopurine 20 mg/ml oral susp</i> | 3 | CC | PA |
| <i>mercaptopurine 50 mg tablet</i> | 1 | CC | |
| <i>methotrexate sodium 2.5 mg tablet</i> | 1 | | |
| <i>methotrexate sodium 25 mg/ml vial</i> | 1 | QL | 16 / 28 days |
| <i>methotrexate sodium/pf 25 mg/ml vial</i> | 1 | QL CC | 16 / 28 days |
| MYLERAN | 3 | CC | PA |
| NERLYNX | 3 | CC | PA |
| <i>nilotinib hcl</i> | 3 | CC | PA |
| <i>nilotinib tartrate</i> | 3 | CC | PA |
| NINLARO | 3 | CC | PA |
| NUBEQA | 3 | CC | PA |
| ODOMZO | 3 | CC | PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| OGSIVEO | 3 | CC PA |
| OJJAARA | 3 | CC PA |
| ONUREG | 3 | CC PA |
| ORSERDU | 3 | CC PA |
| <i>pazopanib hcl</i> | 3 | CC PA |
| PEMAZYRE | 3 | CC PA |
| PIQRAY | 3 | CC PA |
| POMALYST | 3 | CC PA |
| QINLOCK | 3 | CC PA |
| RETEVMO (40 MG CAPSULE, 40 MG TABLET, 80 MG CAPSULE, 80 MG TABLET, 120 MG TABLET, 160 MG TABLET) | 3 | CC PA |
| REVUFORJ | 3 | CC PA |
| REZLIDHIA | 3 | CC PA |
| ROMVIMZA | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|----|
| ROZLYTREK (100 MG CAPSULE, 200 MG CAPSULE) | 3 | CC | PA |
| RUBRACA | 3 | CC | PA |
| RYDAPT | 3 | CC | PA |
| SCEMBLIX | 3 | CC | PA |
| <i>sorafenib tosylate</i> | 3 | CC | PA |
| SPRYCEL | 3 | CC | PA |
| STIVARGA | 3 | CC | PA |
| <i>sunitinib malate</i> | 3 | CC | PA |
| SYNRIBO | 3 | CC | PA |
| TABLOID | 3 | PA | |
| TABRECTA | 3 | CC | PA |
| TAFINLAR (50 MG CAPSULE, 75 MG CAPSULE) | 3 | CC | PA |
| TAGRISSO | 3 | CC | PA |
| TALZENNA | 3 | CC | PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| TAZVERIK | 3 | CC PA |
| <i>temozolomide</i> | 3 | CC PA |
| TEPMETKO | 3 | CC PA |
| TIBSOVO | 3 | CC PA |
| <i>tretinoin 10 mg capsule</i> | 3 | CC PA |
| TRUQAP | 3 | CC PA |
| TUKYSA | 3 | CC PA |
| TURALIO | 3 | CC PA |
| VANFLYTA | 3 | CC PA |
| VENCLEXTA | 3 | CC PA |
| VENCLEXTA STARTING PACK | 3 | CC PA |
| VERZENIO | 3 | CC PA |
| VITRAKVI (20 MG/ML SOLUTION, 25 MG CAPSULE, 100 MG CAPSULE) | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| VIZIMPRO | 3 | CC PA |
| VONJO | 3 | CC PA |
| VORANIGO | 3 | CC PA |
| WELIREG | 3 | CC PA |
| XALKORI (200 MG CAPSULE, 250 MG CAPSULE) | 3 | CC PA |
| XOSPATA | 3 | CC PA |
| XPOVIO | 3 | CC PA |
| XTANDI (40 MG CAPSULE, 40 MG TABLET, 80 MG TABLET) | 3 | CC PA |
| YONSA | 3 | CC PA |
| ZEJULA | 3 | CC PA |
| ZELBORAF | 3 | CC PA |
| ZOLINZA | 3 | CC PA |
| ZYDELIG | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| ZYKADIA | 3 | CC PA |
| ANTIPARKINSONIAN AGENTS (CNS) | | |
| ADAMANTANES (CNS) | | |
| <i>amantadine hcl (50 mg/5 ml solution, 100 mg capsule, 100 mg tablet)</i> | 1 | |
| ANTICHOLINERGIC AGENTS (CNS) | | |
| <i>benztropine mesylate (0.5 mg tablet, 1 mg tablet, 2 mg tablet)</i> | 1 | |
| <i>trihexyphenidyl hcl (2 mg tablet, 2 mg/5 ml solution, 5 mg tablet)</i> | 1 | |
| CATECHOL-O-METHYLTRANSFERASE(COMT)INHIB. | | |
| <i>entacapone</i> | 1 | |
| DOPAMINE PRECURSORS | | |
| <i>carbidopa/levodopa (carbidopa/levodopa 10mg-100mg tab rpd, carbidopa/levodopa 10mg-100mg tablet, carbidopa/levodopa 25mg-100mg tab rpd, carbidopa/levodopa 25mg-100mg tablet, carbidopa/levodopa 25mg-100mg tablet er, carbidopa/levodopa 25mg-250mg tab rpd, carbidopa/levodopa 25mg-250mg tablet, carbidopa/levodopa 50mg-200mg tablet er)</i> | 1 | |
| <i>carbidopa/levodopa/entacapone</i> | 1 | |
| MONOAMINE OXIDASE B INHIBITORS | | |
| <i>selegiline hcl (5 mg capsule, 5 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|----------------|
| ANTIPROTOZOALS | | | |
| ANTIMALARIALS | | | |
| <i>atovaquone/proguanil hcl</i> | 1 | QL | 180 / 365 days |
| <i>chloroquine phosphate (250 mg tablet, 500 mg tablet)</i> | 1 | | |
| <i>hydroxychloroquine sulfate (100 mg tablet, 200 mg tablet, 300 mg tablet, 400 mg tablet)</i> | 1 | | |
| <i>mefloquine hcl</i> | 1 | | |
| <i>primaquine phosphate</i> | 1 | | |
| <i>pyrimethamine 25 mg tablet</i> | 3 | CC PA QPD | 3.0 per day |
| ANTIPROTOZOALS, CRYPTOSPORIDIOSIS | | | |
| <i>nitazoxanide 500 mg tablet</i> | 1 | QL | 30 / 365 days |
| ANTIPROTOZOALS, P JIROVECII PNEUMONIA | | | |
| <i>atovaquone 750 mg/5ml oral susp</i> | 1 | | |
| <i>pentamidine isethionate 300 mg vial-neb</i> | 3 | CC PA QPD | 0.04 per day |
| ANTIPROTOZOALS,NITROIMIDAZOLE-DERIVATIVE | | | |
| <i>tinidazole (250 mg tablet, 500 mg tablet)</i> | 1 | QL | 30 / 365 days |
| ANTIPROTOZOALS,NITROIMIDAZOLE-DERIVATIVE | | | |
| NITROIMIDAZOLE DERIVATIVES, MISC | | | |
| <i>metronidazole (0.75 % gel w/applicator, 250 mg tablet, 500 mg tablet)</i> | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|---------------|
| ANTIPSYCHOTIC AGENTS | | | |
| ATYPICAL ANTIPSYCHOTICS | | | |
| ABILIFY ASIMTUFII 720 MG/2.4ML | 3 | QL CC ST | 2.4/60 days |
| ABILIFY ASIMTUFII 960 MG/3.2ML | 3 | QL CC ST | 3.2 / 60 days |
| ABILIFY MAINTENA (ER 300 MG SYR, ER 300 MG VL, ER 400 MG SYR, ER 400 MG VL) | 3 | QL CC ST | 1 / 30 days |
| <i>aripiprazole (2 mg tablet, 5 mg tablet, 10 mg tablet, 15 mg tablet, 20 mg tablet, 30 mg tablet)</i> | 1 | | |
| ARISTADA ER 1064 MG/3.9 ML SYR | 3 | QL CC ST | 3.9 / 60 days |
| ARISTADA ER 441 MG/1.6 ML SYRN | 3 | QL CC ST | 1.6 / 30 days |
| ARISTADA ER 662 MG/2.4 ML SYRN | 3 | QL CC ST | 2.4 / 30 days |
| ARISTADA ER 882 MG/3.2 ML SYRN | 3 | QL CC ST | 3.2 / 30 days |
| ARISTADA INITIO | 3 | QL CC ST | 2.4 / 42 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|----------------|
| <i>clozapine (25 mg tablet, 50 mg tablet, 100 mg tablet, 200 mg tablet)</i> | 1 | | |
| ERZOFRI 117 MG/0.75 ML SYRINGE | 3 | QL CC ST | 0.75 / 30 days |
| ERZOFRI 156 MG/ML SYRINGE | 3 | QL CC ST | 1 / 30 days |
| ERZOFRI 234 MG/1.5 ML SYRINGE | 3 | QL CC ST | 1.5 / 30 days |
| ERZOFRI 351 MG/2.25 ML SYRINGE | 3 | QL CC ST | 2.25 / 30 days |
| ERZOFRI 39 MG/0.25 ML SYRINGE | 3 | CC ST | |
| ERZOFRI 78 MG/0.5 ML SYRINGE | 3 | QL CC ST | 0.5 / 30 days |
| INVEGA HAFYERA 1,092 MG/3.5 ML | 3 | QL CC ST | 3.5 / 180 days |
| INVEGA HAFYERA 1,560 MG/5 ML | 3 | QL CC ST | 5 / 180 days |
| INVEGA SUSTENNA 117 MG/0.75 ML | 3 | QL CC ST | 0.75 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|----------------|--|
| INVEGA SUSTENNA 156 MG/ML SYRG | 3 | QL | 1 / 30 days | |
| | | CC | | |
| | | ST | | |
| INVEGA SUSTENNA 234 MG/1.5 ML | 3 | QL | 1.5 / 30 days | |
| | | CC | | |
| | | ST | | |
| INVEGA SUSTENNA 39 MG/0.25 ML | 3 | QL | 0.25 / 30 days | |
| | | CC | | |
| | | ST | | |
| INVEGA SUSTENNA 78 MG/0.5 ML | 3 | QL | 0.5 / 30 days | |
| | | CC | | |
| | | ST | | |
| INVEGA TRINZA 273 MG/0.88 ML | 3 | QL | 0.88 / 90 days | |
| | | CC | | |
| | | ST | | |
| INVEGA TRINZA 410 MG/1.32 ML | 3 | QL | 1.32 / 90 days | |
| | | CC | | |
| | | ST | | |
| INVEGA TRINZA 546 MG/1.75 ML | 3 | QL | 1.75 / 90 days | |
| | | CC | | |
| | | ST | | |
| INVEGA TRINZA 819 MG/2.63 ML | 3 | QL | 2.63 / 90 days | |
| | | CC | | |
| | | ST | | |
| <i>olanzapine (2.5 mg tablet, 5 mg tablet, 7.5 mg tablet, 10 mg tablet, 15 mg tablet, 20 mg tablet)</i> | 1 | | | |
| PERSERIS | 3 | QL | 1 / 30 days | |
| | | CC | | |
| | | ST | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|----------------|
| <i>quetiapine fumarate (25 mg tablet, 50 mg tablet, 100 mg tablet, 200 mg tablet, 300 mg tablet, 400 mg tablet)</i> | 1 | | |
| <i>risperidone (0.25 mg tablet, 0.5 mg tablet, 1 mg tablet, 1 mg/ml solution, 2 mg tablet, 3 mg tablet, 4 mg tablet)</i> | 1 | | |
| <i>risperidone microspheres</i> | 3 | QL CC ST | 1 / 14 days |
| UZEDY ER 100 MG/0.28 ML SYRING | 3 | QL CC ST | 0.28 / 30 days |
| UZEDY ER 125 MG/0.35 ML SYRING | 3 | QL CC ST | 0.35 / 30 days |
| UZEDY ER 150 MG/0.42 ML SYRING | 3 | QL CC ST | 0.42 / 30 days |
| UZEDY ER 200 MG/0.56 ML SYRING | 3 | QL CC ST | 0.56 / 30 days |
| UZEDY ER 250 MG/0.7 ML SYRINGE | 3 | QL CC ST | 0.7 / 30 days |
| UZEDY ER 50 MG/0.14 ML SYRINGE | 3 | QL CC ST | 0.14 / 30 days |
| UZEDY ER 75 MG/0.21 ML SYRINGE | 3 | QL CC ST | 0.21 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|-------------|--|
| <i>ziprasidone hcl</i> | 1 | | | |
| ZYPREXA RELPREVV (210 MG VIAL, 210 MG VL KIT, 300 MG VIAL, 300 MG VL KIT) | 3 | QL CC ST | 2 / 30 days | |
| ZYPREXA RELPREVV (405 MG VIAL, 405 MG VL KIT) | 3 | QL CC ST | 1 / 30 days | |
| BUTYROPHENONES | | | | |
| <i>haloperidol (0.5 mg tablet, 1 mg tablet, 2 mg tablet, 5 mg tablet, 10 mg tablet, 20 mg tablet)</i> | 1 | | | |
| <i>haloperidol decanoate (50 mg/ml ampul, 50 mg/ml vial, 100 mg/ml ampul, 100 mg/ml vial)</i> | 3 | CC ST | | |
| <i>haloperidol lactate 2 mg/ml oral conc</i> | 1 | | | |
| DIBENZOXAPINES | | | | |
| <i>loxapine succinate</i> | 1 | | | |
| DIPHENYLBUTYLPERIDINES | | | | |
| <i>pimozide</i> | 1 | | | |
| PHENOTHIAZINES | | | | |
| <i>fluphenazine decanoate 25 mg/ml vial</i> | 3 | QL CC ST | 5 / 30 days | |
| <i>fluphenazine hcl (1 mg tablet, 2.5 mg tablet, 5 mg tablet, 10 mg tablet)</i> | 1 | | | |
| <i>perphenazine (2 mg tablet, 4 mg tablet, 8 mg tablet, 16 mg tablet)</i> | 1 | | | |
| <i>thioridazine hcl (10 mg tablet, 25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| <i>trifluoperazine hcl</i> | 1 | |
| THIOXANTHENES | | |
| <i>thiothixene</i> | 1 | |
| ANTIRETROVIRALS | | |
| ANTIRETROVIRALS, MISCELLANEOUS | | |
| TYBOST | 2 | |
| HIV ENTRY AND FUSION INHIBITORS | | |
| FUZEON | 3 | CC PA |
| <i>maraviroc</i> | 1 | |
| SELZENTRY (25 MG TABLET, 75 MG TABLET) | 2 | |
| HIV INTEGRASE INHIBITOR ANTIRETROVIRALS | | |
| APRETUDE | 2 | HCR \$0 |
| BIKTARVY 50-200-25 MG TABLET | 2 | |
| DOVATO | 2 | |
| ISENTRESS (25 MG TABLET CHEW, 100 MG POWDER PACKET, 100 MG TABLET CHEW, 400 MG TABLET) | 2 | |
| ISENTRESS HD | 2 | |
| JULUCA | 2 | |
| TIVICAY | 2 | |
| VOCABRIA | 2 | |
| HIV NONNUCLEOSIDE REV.TRANScriP. INHIB. | | |
| DELSTRIGO | 2 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| EDURANT | 2 | |
| <i>efavirenz (50 mg capsule, 200 mg capsule, 600 mg tablet)</i> | 1 | |
| <i>efavirenz/lamivudine/tenofovir disoproxil fumarate</i> | 1 | |
| <i>etravirine</i> | 1 | |
| INTELENCE 25 MG TABLET | 2 | |
| <i>nevirapine (50 mg/5 ml oral susp, 100 mg tab er 24h, 200 mg tablet, 400 mg tab er 24h)</i> | 1 | |
| HIV NUCLEOSIDE, NUCLEOTIDE RT INHIBITORS | | |
| <i>abacavir sulfate (20 mg/ml solution, 300 mg tablet)</i> | 1 | |
| <i>abacavir sulfate/lamivudine</i> | 1 | |
| DESCOZY 200-25 MG TABLET | 2 | HCR \$0 |
| <i>didanosine</i> | 1 | |
| <i>efavirenz/emtricitabine/tenofovir disoproxil fumarate</i> | 1 | |
| <i>emtricitabine</i> | 1 | |
| <i>emtricitabine/rilpivirine hcl/tenofovir disoproxil fumarate</i> | 1 | |
| <i>emtricitabine/tenofovir (tdf) 200-300 mg tablet</i> | 1 | HCR \$0 |
| <i>emtricitabine/tenofovir disoproxil fumarate (emtricitabine/tenofovir 100-150 mg tablet, emtricitabine/tenofovir 133-200 mg tablet, emtricitabine/tenofovir 167-250 mg tablet)</i> | 1 | |
| EMTRIVA 10 MG/ML SOLUTION | 2 | |
| GENVOYA | 2 | |
| <i>lamivudine (10 mg/ml solution, 100 mg tablet, 150 mg tablet, 300 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| <i>lamivudine/zidovudine</i> | 1 | |
| ODEFSEY | 2 | |
| <i>stavudine</i> | 1 | |
| STRIBILD | 2 | |
| <i>tenofovir disoproxil fumarate</i> | 1 | |
| TRIUMEQ | 2 | |
| VIREAD (150 MG TABLET, 200 MG TABLET, 250 MG TABLET, POWDER) | 2 | |
| <i>zidovudine (10 mg/ml syrup, 100 mg capsule, 300 mg tablet)</i> | 1 | |
| HIV PROTEASE INHIBITOR ANTIRETROVIRALS | | |
| APTIVUS | 2 | |
| <i>atazanavir sulfate</i> | 1 | |
| <i>darunavir</i> | 1 | |
| <i>darunavir ethanolate</i> | 1 | |
| EVOTAZ | 2 | |
| <i>fosamprenavir calcium</i> | 1 | |
| LEXIVA 50 MG/ML SUSPENSION | 2 | |
| <i>lopinavir/ritonavir (lopinavir/ritonavir 100mg-25mg tablet, lopinavir/ritonavir 200mg-50mg tablet, lopinavir/ritonavir 400-100/5 solution)</i> | 1 | |
| PREZCOBIX 800 MG-150 MG TABLET | 2 | |
| PREZISTA (75 MG TABLET, 100 MG/ML SUSPENSION, 150 MG TABLET) | 2 | |
| REYATAZ 50 MG POWDER PACKET | 2 | |
| <i>ritonavir</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| SYMTUZA | 3 | CC | PA |
| VIRACEPT | 2 | | |
| ANTITHROMBOTIC AGENTS | | | |
| PLATELET-AGGREGATION INHIBITORS | | | |
| <i>cilostazol</i> | 1 | QL | 60 / 30 days |
| <i>clopidogrel bisulfate 75 mg tablet</i> | 1 | | |
| <i>dipyridamole (25 mg tablet, 50 mg tablet, 75 mg tablet)</i> | 1 | | |
| <i>prasugrel hcl</i> | 1 | QL | 30 / 30 days |
| <i>ticagrelor</i> | 1 | QL | 60 / 30 days |
| PLATELET-REDUCING AGENTS | | | |
| <i>anagrelide hcl</i> | 1 | | |
| ANTITOXINS, IMMUNE GLOB, TOXOIDS, VACCINES | | | |
| TOXOIDS | | | |
| ADACEL TDAP (SYRINGE, VIAL) | 2 | \$0 | |
| BOOSTRIX TDAP VACCINE SYRINGE | 2 | \$0 | |
| TENIVAC (SYRINGE, VIAL) | 2 | \$0 | |
| <i>tetanus and diphtheria toxoids, adult</i> | 2 | \$0 | |
| VACCINES | | | |
| ABRYSVO | 2 | \$0 | |
| AFLURIA 2025-2026 | 2 | \$0 | |
| AFLURIA 2025-2026 (3YR UP) | 2 | \$0 | |
| AFLURIA TRIV 2024-25 (3YR UP) | 2 | QL \$0 | 1 fill/year |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|---------------------|
| AFLURIA TRIVALENT 2024-25 | 2 | QL \$0 | 1 fill/year |
| AREXVY | 2 | AL \$0 | At least 50 yrs old |
| BEXSERO | 2 | \$0 | |
| CAPVAXIVE | 2 | \$0 | |
| COMIRNATY 2024-2025 | 2 | \$0 | |
| ENGERIX-B ADULT (20 MCG/ML SYRN, 20 MCG/ML VIAL) | 2 | \$0 | |
| FLUAD 2025-2026 | 2 | \$0 | |
| FLUARIX 2025-2026 | 2 | \$0 | |
| FLUARIX TRIVALENT 2024-2025 | 2 | QL \$0 | 1 fill/year |
| FLUBLOK 2025-2026 | 2 | \$0 | |
| FLUBLOK TRIVALENT 2024-2025 | 2 | QL \$0 | 1 fill/year |
| FLUCELVAX 2025-2026 (2025-2026 SYRINGE, 2025-2026 VIAL) | 2 | \$0 | |
| FLUCELVAX TRIVALENT 2024-2025 (2024-2025 SYR, 2024-2025 VL) | 2 | QL \$0 | 1 fill/year |
| FLULAVAL 2025-2026 | 2 | \$0 | |
| FLULAVAL TRIVALENT 2024-2025 | 2 | QL \$0 | 1 fill/year |
| FLUMIST 2025-2026 | 2 | \$0 | |
| FLUMIST HOME 2025-2026 | 2 | \$0 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|----------------------------------|
| FLUZONE 2025-2026 (2025-2026 SYRINGE, 2025-2026 VIAL) | 2 | \$0 |
| FLUZONE HIGH-DOSE 2025-2026 | 2 | \$0 |
| FLUZONE TRIV SOUTHERN HEM 2025 | 2 | QL \$0 1 fill/year |
| FLUZONE TRIVALENT 2024-2025 (2024-25 SYRG, 2024-25 VIAL) | 2 | QL \$0 1 fill/year |
| GARDASIL 9 (9 SYRINGE, 9 VIAL) | 2 | \$0 |
| HAVRIX | 2 | QL \$0 1 fill/6 months |
| HEPLISAV-B | 2 | \$0 |
| IMOVAX RABIES VACCINE | 2 | \$0 |
| IPOP (SINGLE DOSE SYRINGE, VIAL) | 2 | |
| IXCHIQ | 2 | \$0 |
| IXIARO | 2 | \$0 |
| JYNNEOS | 2 | \$0 |
| JYNNEOS (NATIONAL STOCKPILE) | 2 | \$0 |
| M-M-R II VACCINE | 2 | \$0 |
| MENQUADFI | 2 | \$0 |
| MENVEO A-C-Y-W-135-DIP (1 VIAL-A-C-Y-W-135-DIP, A-C-Y-W KIT (2 VIALS)) | 2 | \$0 |
| MRESVIA | 2 | AL \$0 At least 60 yrs old |
| NOVAVAX COVID 2024-2025 (EUA) | 2 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|------------------------------|
| PENBRAYA | 2 | \$0 |
| PENMENVY MEN A-B-C-W-Y | 2 | \$0 |
| PENMENVY MENACWY COMPONENT | 2 | \$0 |
| PENMENVY MENB COMPONENT | 2 | \$0 |
| PNEUMOVAX 23 (23 SYRINGE, 23 VIAL) | 2 | \$0 |
| PREVNAR 13 | 2 | \$0 |
| PREVNAR 20 | 2 | \$0 |
| PRIORIX | 2 | \$0 |
| RABAVERT | 2 | \$0 |
| RECOMBIVAX HB (5 MCG/0.5 ML SYR, 5 MCG/0.5 ML VL, 10 MCG/ML SYR, 10 MCG/ML VIAL, 40 MCG/ML VIAL) | 2 | \$0 |
| SHINGRIX | 2 | \$0 |
| SPIKEVAX 2024-2025 | 2 | \$0 |
| STAMARIL | 2 | \$0 |
| TICOVAC | 2 | \$0 |
| TRUMENBA | 2 | \$0 |
| TWINRIX | 2 | \$0 |
| TYPHIM VI (25 MCG/0.5 ML AL, 25 MCG/0.5 ML SYRNG) | 2 | \$0 |
| VAQTA (25 UNITS/0.5 ML SYRINGE, 25 UNITS/0.5 ML VIAL, 50 UNITS/ML SYRINGE, 50 UNITS/ML VIAL) | 2 | QL \$0 1 fill/6 months |
| VARIVAX VACCINE | 2 | \$0 |
| VAXCHORA VACCINE | 2 | \$0 |
| VAXNEUVANCE | 2 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|------------------------|
| VIVOTIF | 2 | \$0 |
| YF-VAX | 2 | \$0 |
| ANTIULCER AGENTS AND ACID SUPPRESSANTS | | |
| HISTAMINE H2-ANTAGONISTS | | |
| <i>cimetidine (200 mg tablet, 300 mg tablet, 400 mg tablet, 800 mg tablet)</i> | 1 | |
| <i>famotidine (20 mg tablet, 40 mg tablet)</i> | 1 | |
| PROSTAGLANDINS | | |
| <i>misoprostol (100 mcg tablet, 200 mcg tablet)</i> | 1 | \$0 |
| PROTECTANTS | | |
| <i>sucralfate 1 g tablet</i> | 1 | |
| PROTON-PUMP INHIBITORS | | |
| <i>esomeprazole magnesium 20 mg capsule dr</i> | 1 | |
| <i>lansoprazole (15 mg capsule dr, 30 mg capsule dr)</i> | 1 | |
| <i>omeprazole (10 mg capsule dr, 20 mg capsule dr, 40 mg capsule dr)</i> | 1 | |
| <i>pantoprazole sodium (20 mg tablet dr, 40 mg tablet dr)</i> | 1 | |
| <i>rabeprazole sodium 20 mg tablet dr</i> | 1 | |
| ANTIVIRALS (SYSTEMIC) | | |
| ANTIRETROVIRALS | | |
| YEZTUGO (300 MG TABLET, 463.5 MG/1.5 ML VIAL) | 2 | \$0 |
| CORONAVIRUS (COVID-19) | | |
| PAXLOVID | 2 | QPD 6.0 per day \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|------------------|
| ENDONUCLEASE INHIBITORS | | | |
| XOFLUZA | 2 | QL | 2 / 180 days |
| NEURAMINIDASE INHIBITOR ANTIVIRALS | | | |
| <i>oseltamivir phosphate (30 mg capsule, 75 mg capsule)</i> | 1 | | |
| <i>oseltamivir phosphate (6 mg/ml susp recon, 45 mg capsule)</i> | 1 | AL | Up to 12 yrs old |
| RELENZA | 2 | QL | 20 / 30 days |
| NUCLEOSIDE AND NUCLEOTIDE ANTIVIRALS | | | |
| <i>acyclovir (200 mg capsule, 400 mg tablet, 800 mg tablet)</i> | 1 | QL | 150 / 30 days |
| <i>entecavir</i> | 1 | QL | 30 / 30 days |
| <i>famciclovir 125 mg tablet</i> | 1 | QL | 60 / 30 days |
| <i>famciclovir 250 mg tablet</i> | 1 | QL | 90 / 30 days |
| <i>famciclovir 500 mg tablet</i> | 1 | QL | 120 / 30 days |
| LAGEVRIO (EUA) | 2 | QPD \$0 | 8.0 per day |
| <i>ribavirin (200 mg capsule, 200 mg tablet)</i> | 1 | | |
| <i>valacyclovir hcl 1000 mg tablet</i> | 1 | QL | 120 / 30 days |
| <i>valacyclovir hcl 500 mg tablet</i> | 1 | QL | 90 / 30 days |
| <i>valganciclovir hcl 50 mg/ml soln recon</i> | 3 | CC PA QPD | 36.0 per day |
| <i>valganciclovir hcl 450 mg tablet</i> | 3 | CC PA QPD | 4.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------------|
| VEMLIDY | 2 | |
| ANXIOLYTICS, SEDATIVES AND HYPNOTICS | | |
| ANXIOLYTICS, SEDATIVES, AND HYPNOTICS, MISC | | |
| <i>hydroxyzine hcl (10 mg tablet, 25 mg tablet, 50 mg tablet)</i> | 1 | |
| <i>hydroxyzine pamoate (25 mg capsule, 50 mg capsule, 100 mg capsule)</i> | 1 | |
| BARBITURATES (ANXIOLYTIC, SEDATIVE/HYP) | | |
| <i>butalb/acetaminophen/caffeine 50-325-40 tablet</i> | 3 | CC PA QPD 2.0 per day |
| <i>phenobarbital (15 mg tablet, 16.2 mg tablet, 30 mg tablet, 32.4 mg tablet, 60 mg tablet, 64.8 mg tablet, 97.2mg tablet, 100 mg tablet)</i> | 1 | |
| BENZODIAZEPINES (ANXIOLYTIC, SEDATIVE/HYP) | | |
| <i>chlordiazepoxide hcl</i> | 1 | QL 120 / 30 days |
| <i>diazepam (2.5 mg kit, 5-7.5-10mg kit, 12.5-15-20 kit)</i> | 1 | QL 3 / 365 days |
| <i>diazepam (2 mg tablet, 5 mg tablet, 10 mg tablet)</i> | 1 | QL 120 / 30 days |
| <i>lorazepam (0.5 mg tablet, 1 mg tablet)</i> | 1 | QL 90 / 30 days |
| <i>lorazepam 2 mg tablet</i> | 1 | QL 150 / 30 days |
| NAYZILAM | 2 | QL 3 / 365 days |
| <i>temazepam</i> | 1 | QL 30 / 30 days |
| VALTOCO | 2 | QL 3 / 365 days |
| MELATONIN RECEPTOR AGONISTS | | |
| <i>ramelteon</i> | 3 | QL 30 / 30 days CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|---------------|
| NON-BENZODIAZEPINE ANXIOLYTICS | | | |
| <i>buspirone hcl (5 mg tablet, 7.5 mg tablet, 10 mg tablet, 15 mg tablet, 30 mg tablet)</i> | 1 | | |
| NON-BENZODIAZEPINE HYPNOTICS | | | |
| <i>eszopiclone (2 mg tablet, 3 mg tablet)</i> | 3 | QL CC ST | 30 / 30 days |
| <i>eszopiclone 1 mg tablet</i> | 1 | QL | 30 / 30 days |
| <i>zaleplon</i> | 1 | QL | 30 / 30 days |
| <i>zolpidem tartrate 10 mg tablet</i> | 1 | QL ST | 30 / 30 days |
| <i>zolpidem tartrate 5 mg tablet</i> | 1 | QL | 30 / 30 days |
| <i>zolpidem tartrate (6.25 mg tab mphase, 12.5 mg tab mphase)</i> | 3 | QL CC ST | 30 / 30 days |
| AUTONOMIC DRUGS | | | |
| PARASYMPATHOMIMETIC (CHOLINERGIC AGENTS) | | | |
| <i>bethanechol chloride (5 mg tablet, 10 mg tablet, 25 mg tablet, 50 mg tablet)</i> | 1 | | |
| <i>cevimeline hcl</i> | 1 | QL | 90 / 30 days |
| <i>donepezil hcl (5 mg tab rapdis, 5 mg tablet, 10 mg tab rapdis, 10 mg tablet, 23 mg tablet)</i> | 1 | | |
| <i>galantamine hbr (4 mg tablet, 4 mg/ml solution, 8 mg cap24h pel, 8 mg tablet, 12 mg tablet, 16 mg cap24h pel, 24 mg cap24h pel)</i> | 1 | | |
| <i>pilocarpine hcl 5 mg tablet</i> | 1 | QL | 180 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|---------------|
| <i>pilocarpine hcl 7.5 mg tablet</i> | 1 | QL | 120 / 30 days |
| <i>pyridostigmine bromide (60 mg tablet, 60 mg/5 ml solution, 180 mg tablet er)</i> | 1 | | |
| <i>rivastigmine</i> | 1 | | |
| <i>rivastigmine tartrate</i> | 1 | | |

SMOKING CESSATION AGENTS

| | | | |
|--|---|------------------------|--------------|
| NICOTROL | 3 | CC PA QPD \$0 | 16.8 per day |
| NICOTROL NS | 3 | CC PA QPD \$0 | 4.0 per day |
| <i>varenicline tartrate 0.5 (11)-1 tab ds pk</i> | 1 | QL \$0 | 53 / 28 days |
| <i>varenicline tartrate (0.5 mg tablet, 1 mg tablet)</i> | 1 | \$0 | |

BETA-ADRENERGIC AGONISTS

SELECTIVE BETA-2-ADRENERGIC AGONISTS

| | | | |
|--|---|-----|--------------|
| <i>albuterol sulfate 90 mcg hfa aer ad</i> | 1 | QPD | 1.2 per day |
| <i>albuterol sulfate 5 mg/ml solution</i> | 1 | QPD | 3.4 per day |
| <i>albuterol sulfate (2 mg tablet, 4 mg tablet)</i> | 1 | QPD | 4.0 per day |
| <i>albuterol sulfate (0.63mg/3ml vial-neb, 1.25mg/3ml vial-neb, 2.5 mg/0.5 vial-neb)</i> | 1 | | |
| <i>albuterol sulfate 2.5 mg/3ml vial-neb</i> | 1 | QPD | 18.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| BREYNA | 1 | QPD | 0.7 per day |
| <i>budesonide/formoterol fumarate</i> | 1 | QPD | 0.7 per day |
| <i>fluticasone propionate/salmeterol xinafoate (propion/salmeterol 55-14 mcg aer pow ba, propion/salmeterol 113-14 mcg aer pow ba, propion/salmeterol 232-14 mcg aer pow ba)</i> | 1 | QPD | 0.04 per day |
| <i>fluticasone propionate/salmeterol xinafoate (propion/salmeterol 100-50 mcg blst w/dev, propion/salmeterol 250-50 mcg blst w/dev, propion/salmeterol 500-50 mcg blst w/dev)</i> | 1 | QPD | 2.0 per day |
| <i>levalbuterol hcl (0.63mg/3ml vial-neb, 1.25mg/3ml vial-neb)</i> | 3 | ST QPD | 12.0 per day |
| <i>levalbuterol hcl 0.31mg/3ml vial-neb</i> | 3 | ST QPD | 8.0 per day |
| <i>levalbuterol hcl 1.25mg/0.5 vial-neb</i> | 3 | ST QPD | 5.0 per day |
| <i>levalbuterol tartrate</i> | 3 | ST QPD | 1.0 per day |
| STRIVERDI RESPIMAT | 2 | QPD | 0.15 per day |
| <i>terbutaline sulfate (2.5 mg tablet, 5 mg tablet)</i> | 1 | | |
| WIXELA INHUB | 1 | QPD | 2.0 per day |
| BLOOD FORMATION, COAGULATION, THROMBOSIS | | | |
| BLOOD FORM.,COAG,THROMBOSIS AGENTS MISC. | | | |
| OXBRYTA 500 MG TABLET | 3 | CC PA | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| HEMATOPOIETIC AGENTS | | | |
| <i>eltrombopag olamine (12.5 mg tablet, 25 mg tablet, 50 mg tablet, 75 mg tablet)</i> | 3 | CC PA QPD | 1.0 per day |
| FYLNETRA | 3 | CC PA QPD | 0.08 per day |
| JESDUVROQ | 3 | CC PA | |
| <i>plerixafor</i> | 3 | CC PA | |
| RELEUKO 300 MCG/0.5 ML SYRINGE | 3 | CC PA QPD | 0.25 per day |
| RELEUKO 480 MCG/0.8 ML SYRINGE | 3 | CC PA QPD | 0.4 per day |
| RELEUKO 300 MCG/ML VIAL | 3 | CC PA QPD | 0.5 per day |
| RELEUKO 480 MCG/1.6 ML VIAL | 3 | CC PA QPD | 0.8 per day |
| RETACRIT (2,000 UNIT/ML VIAL, 3,000 UNIT/ML VIAL, 4,000 UNIT/ML VIAL, 10,000 UNIT/ML VIAL, 20,000 UNIT/ML VIAL, 40,000 UNIT/ML VIAL) | 3 | CC PA QPD | 0.43 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| RETACRIT 20,000 UNIT/2 ML VIAL | 3 | CC PA QPD | 0.86 per day |
| HEMORRHEOLOGIC AGENTS | | | |
| <i>pentoxifylline 400 mg tablet er</i> | 1 | QL | 90 / 30 days |
| CALCINEURIN INHIBITORS (90:28) | | | |
| CALCINEURIN INHIBITORS, MISC (90:28) | | | |
| <i>cyclosporine, modified (25 mg capsule, 50 mg capsule, 100 mg capsule)</i> | 1 | | |
| <i>tacrolimus (0.5 mg capsule, 1 mg capsule, 5 mg capsule)</i> | 1 | | |
| CALCIUM-CHANNEL BLOCKING AGENTS | | | |
| DIHYDROPYRIDINES | | | |
| <i>amlodipine besylate (2.5 mg tablet, 5 mg tablet, 10 mg tablet)</i> | 1 | | |
| <i>amlodipine besylate/benazepril hcl</i> | 1 | | |
| <i>amlodipine besylate/valsartan</i> | 1 | | |
| <i>felodipine</i> | 1 | | |
| <i>nifedipine (30 mg tab er 24, 30 mg tablet er, 60 mg tab er 24, 60 mg tablet er, 90 mg tab er 24, 90 mg tablet er)</i> | 1 | | |
| CARDIAC DRUGS | | | |
| CARDIAC DRUGS, MISCELLANEOUS | | | |
| <i>ranolazine</i> | 3 | ST | |
| CARDIOTONIC AGENTS | | | |
| DIGITEK | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| <i>digoxin (50 mcg/ml solution, 125 mcg tablet, 250 mcg tablet)</i> | 1 | |
| CARDIOVASCULAR DRUGS | | |
| ALPHA-ADRENERGIC BLOCKING AGENTS | | |
| <i>doxazosin mesylate (1 mg tablet, 2 mg tablet, 4 mg tablet, 8 mg tablet)</i> | 1 | |
| <i>prazosin hcl (1 mg capsule, 2 mg capsule, 5 mg capsule)</i> | 1 | |
| <i>terazosin hcl</i> | 1 | |
| BETA-ADRENERGIC BLOCKING AGENTS | | |
| <i>atenolol (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| <i>atenolol/chlorthalidone</i> | 1 | |
| <i>bisoprolol fumarate (5 mg tablet, 10 mg tablet)</i> | 1 | |
| <i>bisoprolol fumarate/hydrochlorothiazide</i> | 1 | |
| <i>carvedilol</i> | 1 | |
| <i>labetalol hcl (100 mg tablet, 200 mg tablet, 300 mg tablet)</i> | 1 | |
| <i>metoprolol succinate</i> | 1 | |
| <i>metoprolol tartrate (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| <i>propranolol hcl (10 mg tablet, 20 mg tablet, 40 mg tablet, 60 mg cap sa 24h, 60 mg tablet, 80 mg cap sa 24h, 80 mg tablet, 120 mg cap sa 24h, 160 mg cap sa 24h)</i> | 1 | |
| <i>SOTALOL AF</i> | 1 | |
| <i>sotalol hcl (80 mg tablet, 120 mg tablet, 160 mg tablet, 240 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| CARDIOVASCULAR DRUGS, NSAID ANTI-INFL | | |
| <i>colchicine (0.6 mg capsule, 0.6 mg tablet)</i> | 1 | QL 60 / 30 days |
| CENTRAL ALPHA-AGONISTS | | |
| <i>clonidine (0.2mg/24hr patch tdwk, 0.3mg/24hr patch tdwk)</i> | 1 | QL 8 / 28 days |
| <i>clonidine 0.1mg/24hr patch tdwk</i> | 1 | QL 4 / 28 days |
| <i>clonidine hcl (0.1 mg tab er 12h, 0.1 mg tablet, 0.2 mg tablet, 0.3 mg tablet)</i> | 1 | |
| <i>guanfacine hcl (1 mg tab er 24h, 1 mg tablet, 2 mg tab er 24h, 2 mg tablet, 3 mg tab er 24h, 4 mg tab er 24h)</i> | 1 | |
| <i>methyldopa</i> | 1 | |
| CENTRAL NERVOUS SYSTEM AGENTS | | |
| AMYOTROPHIC LATERAL SCLEROSIS(ALS) AGENT | | |
| <i>riluzole</i> | 1 | |
| ANTIMANIC AGENTS | | |
| <i>lithium carbonate (150 mg capsule, 300 mg capsule, 300 mg tablet, 300 mg tablet er, 450 mg tablet er, 600 mg capsule)</i> | 1 | |
| CENTRAL NERVOUS SYSTEM AGENTS, MISC. | | |
| <i>carbidopa 25 mg tablet</i> | 1 | |
| <i>memantine hcl (2 mg/ml solution, 5 mg tablet, 5 mg-10 mg tab ds pk, 7 mg cap spr 24, 10 mg tablet, 14 mg cap spr 24, 21 mg cap spr 24, 28 mg cap spr 24)</i> | 1 | |
| <i>memantine hcl/donepezil hcl</i> | 1 | |
| NAMENDA XR TITRATION PACK | 2 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|---|--|
| NAMZARIC (7 MG-10 MG CAPSULE, TITRATION PACK) | 2 | | |
| NUEDEXTA | 3 | CC PA QPD 2.0 per day | |
| OPIOID ANTAGONISTS (28:10) | | | |
| KLOXXADO | 2 | | |
| <i>naloxone hcl (0.4 mg/ml cartridge, 0.4 mg/ml syringe, 0.4 mg/ml vial, 1 mg/ml syringe, 4 mg spray)</i> | 1 | | |
| <i>naltrexone hcl 50 mg tablet</i> | 1 | | |
| OPVEE | 2 | | |
| VIVITROL | 2 | | |
| VESICULAR MONOAMINE TRANSPORT2 INHIBITOR | | | |
| AUSTEDO | 3 | QL 120 / 30 days CC PA QPD 4.0 per day | |
| INGREZZA | 3 | CC PA QPD 1.0 per day | |
| INGREZZA INITIATION PK(TARDIV) | 3 | CC PA QPD 1.0 per day | |
| <i>tetrabenazine</i> | 3 | CC PA QPD 4.0 per day | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------------|
| CEPHALOSPORIN ANTIBIOTICS | | |
| 1ST GENERATION CEPHALOSPORIN ANTIBIOTICS | | |
| <i>cefadroxil 500 mg capsule</i> | 1 | |
| <i>cephalexin (250 mg capsule, 500 mg capsule)</i> | 1 | |
| 2ND GENERATION CEPHALOSPORIN ANTIBIOTICS | | |
| <i>cefaclor (250 mg capsule, 500 mg capsule)</i> | 1 | |
| <i>cefprozil 125 mg/5ml susp recon</i> | 1 | AL Up to 12 yrs old |
| <i>cefprozil (250 mg tablet, 500 mg tablet)</i> | 1 | |
| <i>cefuroxime axetil</i> | 1 | |
| 3RD GENERATION CEPHALOSPORIN ANTIBIOTICS | | |
| <i>cefdinir 300 mg capsule</i> | 1 | |
| <i>cefixime 400 mg capsule</i> | 1 | |
| <i>cefpodoxime proxetil (100 mg tablet, 200 mg tablet)</i> | 1 | |
| COMPLEMENT INHIBITORS (92:32) | | |
| BRADYKININ RECEPTOR ANTAGONISTS | | |
| <i>icatibant acetate</i> | 3 | CC PA QPD 3.0 per day |
| CONSTIPATION THERAPY | | |
| CHLORIDE CHANNEL ACTIVATORS | | |
| <i>lubiprostone</i> | 3 | CC PA QPD 2.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|--|
| GUANYLATE CYCLASE C (GCC) RECEPT AGONIST | | |
| LINZESS | 2 | |
| OPIOID ANTAGONISTS (56:18) | | |
| MOVANTIK | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 1.0 per day </div> |
| SYMPROIC | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 1.0 per day </div> |
| CYSTIC FIBROSIS (CFTR) MODULATORS | | |
| CYSTIC FIBROSIS (CFTR) CORRECTORS | | |
| ORKAMBI (100-125 MG GRANULE PKT, 150-188 MG GRANULE PKT) | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 2.0 per day </div> |
| ORKAMBI (100 MG TABLET, 200 MG TABLET) | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 4.0 per day </div> |
| SYMDEKO | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 2.0 per day </div> |
| TRIKAFTA 100-50-75 MG/150 MG | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 3.0 per day </div> |
| CYSTIC FIBROSIS (CFTR) POTENTIATORS | | |
| KALYDECO (25 MG GRANULES PACKET, 50 MG GRANULES PACKET, 75 MG GRANULES PACKET, 150 MG TABLET) | 3 | <div style="display: flex; justify-content: space-between;"> CC PA QPD 2.0 per day </div> |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| DENTAL AGENTS | | | |
| NUTRITIONAL SUPPLEMENTS | | | |
| DENTA 5000 PLUS | 1 | | |
| DENTAGEL | 1 | | |
| <i>fluoride (sodium) (1.1 % cream (g), 1.1 % gel (gram))</i> | 1 | | |
| FRAICHE 5000 | 1 | | |
| SF | 1 | | |
| SF 5000 PLUS | 1 | | |
| SODIUM FLUORIDE 5000 PLUS | 1 | | |
| DEPIGMENTING AND PIGMENTING AGENTS | | | |
| PIGMENTING AGENTS | | | |
| <i>methoxsalen 10 mg cap lg rap</i> | 1 | QL | 12 / 28 days |
| DEVICES | | | |
| <i>blood-glucose meter,continuous</i> | 3 | QL CC PA | 1 / 365 days |
| <i>blood-glucose sensor</i> | 3 | QL CC PA | 3 / 30 days |
| <i>compressor, for nebulizer</i> | 2 | QL | 2 / 365 days |
| <i>diabetic needles</i> | 1 | | |
| <i>diabetic syringes</i> | 1 | | |
| <i>flash glucose scanning reader</i> | 3 | QL CC PA | 1 / 365 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| <i>flash glucose sensor</i> | 3 | QL CC PA | 3 / 30 days |
| <i>inhaler, assist devices</i> | 2 | QL | 2 / 365 days |
| <i>inhaler, assist device, accessory each</i> | 2 | QL | 2 / 365 days |
| <i>inhaler, assist device with large mask</i> | 2 | QL | 2 / 365 days |
| <i>inhaler, assist device with medium mask</i> | 2 | QL | 2 / 365 days |
| <i>inhaler, assist device with small mask</i> | 2 | QL | 2 / 365 days |
| <i>mucus clearing device</i> | 2 | QL | 2 / 365 days |
| <i>nasal exhalation resistance dev each</i> | 2 | QL | 2 / 365 days |
| <i>nebulizer</i> | 2 | QL | 2 / 365 days |
| <i>nebulizer and compressor</i> | 2 | QL | 2 / 365 days |
| <i>peak flow meter</i> | 2 | QL | 2 / 365 days |
| <i>peak flow meter/inhaler, assist devices</i> | 2 | QL | 2 / 365 days |
| <i>spirometers and accessories</i> | 2 | QL | 2 / 365 days |

DISEASE-MODIFYING ANTIRHEUMATIC DRUGS

DISEASE-MODIFYING ANTIRHEUMAT DRUGS MISC

| | | | |
|---|---|-----------------|--------------|
| ORENCIA (50 MG/0.4 ML SYRINGE, 87.5 MG/0.7 ML SYRINGE, 125 MG/ML SYRINGE) | 3 | CC PA QPD | 0.15 per day |
| ORENCIA CLICKJECT | 3 | CC PA QPD | 0.15 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| MONOCARBOXYLIC ACID AMIDE AGENTS | | |
| <i>leflunomide (10 mg tablet, 20 mg tablet)</i> | 1 | |
| DIURETICS | | |
| LOOP DIURETICS (40:28) | | |
| <i>bumetanide (0.5 mg tablet, 1 mg tablet, 2 mg tablet)</i> | 1 | |
| <i>furosemide (10 mg/ml solution, 20 mg tablet, 40 mg tablet, 40mg/5ml solution, 80 mg tablet)</i> | 1 | |
| <i>torsemide</i> | 1 | |
| OSMOTIC DIURETICS | | |
| <i>urea 40 % cream (g)</i> | 1 | QL 198.4 / 30 days |
| POTASSIUM-SPARING DIURETICS | | |
| <i>amiloride hcl</i> | 1 | |
| <i>triamterene/hydrochlorothiazide (triamterene/hydrochlorothiazid 37.5-25 mg capsule, triamterene/hydrochlorothiazid 37.5-25 mg tablet, triamterene/hydrochlorothiazid 75 mg-50mg tablet)</i> | 1 | |
| THIAZIDE DIURETICS | | |
| <i>hydrochlorothiazide (12.5 mg capsule, 12.5 mg tablet, 25 mg tablet, 50 mg tablet)</i> | 1 | |
| THIAZIDE-LIKE DIURETICS | | |
| <i>chlorthalidone</i> | 1 | |
| <i>indapamide</i> | 1 | |
| <i>metolazone</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| DOPAMINE RECEPTOR AGONISTS | | |
| ERGOT-DERIV. DOPAMINE RECEPTOR AGONISTS | | |
| <i>bromocriptine mesylate (2.5 mg tablet, 5 mg capsule)</i> | 1 | |
| <i>cabergoline</i> | 1 | |
| NONERGOT-DERIV.DOPAMINE RECEPTOR AGONIST | | |
| <i>pramipexole di-hcl (0.125 mg tablet, 0.25 mg tablet, 0.5 mg tablet, 0.75 mg tablet, 1 mg tablet, 1.5 mg tablet)</i> | 1 | |
| <i>ropinirole hcl (0.25 mg tablet, 0.5 mg tablet, 1 mg tablet, 2 mg tablet, 3 mg tablet, 4 mg tablet, 5 mg tablet)</i> | 1 | |
| ELECTROLYTIC, CALORIC, AND WATER BALANCE | | |
| ALKALINIZING AGENTS | | |
| <i>potassium citrate</i> | 1 | QL 120 / 30 days |
| AMMONIA DETOXICANTS | | |
| CONSTULOSE | 1 | |
| ENULOSE | 1 | |
| GENERLAC | 1 | |
| <i>lactulose</i> | 1 | |
| <i>sodium phenylbutyrate 0.94 g/g powder</i> | 1 | |
| IRRIGATING SOLUTIONS | | |
| <i>sodium chloride for inhalation (0.9 % vial-neb, 3 % vial-neb, 7 % vial-neb)</i> | 1 | |
| REPLACEMENT PREPARATIONS | | |
| KLOR-CON M10 | 1 | |
| KLOR-CON M20 | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| <i>potassium chloride (8 capsule er, 8 tablet er, 10 capsule er, 10 tab er prt, 10 tablet er, 20 tab er prt, 20 tablet er)</i> | 1 | | |
| URICOSURIC AGENTS | | | |
| <i>probenecid</i> | 1 | | |
| EMOLLIENTS, DEMULCENTS, AND PROTECTANTS | | | |
| BASIC LOTIONS AND LINIMENTS | | | |
| <i>ammonium lactate 12 % lotion</i> | 1 | | |
| BASIC OINTMENTS AND PROTECTANTS | | | |
| <i>ammonium lactate 12 % cream (g)</i> | 1 | | |
| <i>calcipotriene (0.005 % cream (g), 0.005 % oint. (g), 0.005 % solution)</i> | 1 | QL | 60 / 30 days |
| <i>nitroglycerin 0.4% (w/w) oint. (g)</i> | 3 | QL CC PA | 30 / 30 days |
| SANTYL | 2 | QL | 30 / 30 days |
| ENZYMES | | | |
| ENZYME COFACTORS/CHAPERONES | | | |
| <i>nitisinone (2 mg capsule, 5 mg capsule, 10 mg capsule)</i> | 3 | CC PA | |
| <i>nitisinone 20 mg capsule</i> | 3 | CC PA | |
| ENZYME INHIBITORS | | | |
| CERDELGA | 3 | CC PA QPD | 2.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------------|--|--|
| <i>miglustat</i> | 3 | CC PA QPD 3.0 per day | | |
| OPFOLDA | 3 | CC PA QPD 0.3 per day | | |
| YARGESA | 3 | CC PA QPD 3.0 per day | | |
| ESTROGENS AND ANTIESTROGENS | | | | |
| ESTROGEN AGONIST-ANTAGONISTS | | | | |
| CLOMID | 3 | CC PA | | |
| <i>clomiphene citrate 50 mg tablet</i> | 3 | CC PA | | |
| <i>raloxifene hcl</i> | 1 | \$0 | | |
| SOLTAMOX | 3 | CC PA | | |
| <i>tamoxifen citrate (10 mg tablet, 20 mg tablet)</i> | 1 | \$0 | | |
| <i>toremifene citrate</i> | 3 | CC PA | | |
| ESTROGENS | | | | |
| CLIMARA PRO | 2 | QL 4 / 28 days | | |
| COMBIPATCH | 2 | QL 8 / 28 days | | |
| DEPO-ESTRADIOL | 2 | | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| DOTTI (0.0375 MG PATCH, 0.05 MG PATCH, 0.075 MG PATCH, 0.1 MG PATCH) | 1 | QL | 24 / 84 days |
| DOTTI 0.025 MG PATCH | 1 | | |
| <i>estradiol 1.25 g gel md pmp</i> | 1 | QL | 50 / 30 days |
| <i>estradiol (.0375mg/24 patch tds, 0.05mg/24h patch tds, .075mg/24h patch tds, 0.1mg/24hr patch tds)</i> | 1 | QL | 24 / 84 days |
| <i>estradiol (.025mg/24h patch tdwk, .0375mg/24 patch tdwk, 0.05mg/24h patch tdwk, 0.06mg/24h patch tdwk, .075mg/24h patch tdwk, 0.1mg/24hr patch tdwk)</i> | 1 | QL | 4 / 28 days |
| <i>estradiol (0.01 % cream/appl, .025mg/24h patch tds, 0.5 mg tablet, 1 mg tablet, 2 mg tablet, 10 mcg tablet)</i> | 1 | | |
| <i>estradiol valerate (10 mg/ml vial, 20 mg/ml vial, 40 mg/ml vial)</i> | 1 | QL | 10 / 90 days |
| ESTRING | 3 | QL ST | 1 / 90 days |
| FYAVOLV 1 MG-5 MCG TABLET | 1 | | |
| JINTELI | 1 | | |
| LYLLANA (0.0375 MG PATCH, 0.05 MG PATCH, 0.075 MG PATCH, 0.1 MG PATCH) | 1 | QL | 24 / 84 days |
| LYLLANA 0.025 MG PATCH | 1 | | |
| MENOSTAR | 2 | | |
| <i>norethindrone ac-eth estradiol 1mg-5mcg tablet</i> | 1 | | |
| PREMARIN VAGINAL CREAM-APPL | 3 | ST | |
| PREMARIN (0.3 MG TABLET, 0.45 MG TABLET, 0.625 MG TABLET, 0.9 MG TABLET, 1.25 MG TABLET) | 2 | | |
| PREMPHASE | 2 | | |
| PREMPRO | 2 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|--------------|--|
| YUVAFEM | 1 | | | |
| EYE, EAR, NOSE AND THROAT (EENT) PREPS. | | | | |
| ANTI-INFLAMMATORY AGENTS (EENT) | | | | |
| <i>cyclosporine 0.05 % droperette</i> | 3 | CC PA QPD | 2.0 per day | |
| ANTIALLERGIC AGENTS | | | | |
| <i>azelastine hcl 0.05 % drops</i> | 1 | | | |
| <i>azelastine hcl 137 mcg spray/pump</i> | 1 | QL | 30 / 30 days | |
| <i>cromolyn sodium 4 % drops</i> | 1 | | | |
| <i>olopatadine hcl (0.1 % drops, 0.2 % drops)</i> | 3 | CC ST QPD | 0.17 per day | |
| EENT DRUGS, MISCELLANEOUS | | | | |
| <i>ipratropium bromide (21 mcg spray, 42 mcg spray)</i> | 1 | | | |
| LOCAL ANESTHETICS (EENT) | | | | |
| <i>lidocaine hcl 2 % solution</i> | 1 | | | |
| <i>proparacaine hcl 0.5 % drops</i> | 1 | | | |
| MYDRIATICS | | | | |
| <i>atropine sulfate 1 % drops</i> | 1 | | | |
| <i>cyclopentolate hcl</i> | 1 | | | |
| <i>tropicamide 1 % drops</i> | 1 | | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------------|
| FIRST GENERATION ANTIHISTAMINES | | |
| FIRST GEN. ANTIHIST. DERIVATIVES, MISC. | | |
| <i>cyproheptadine hcl 4 mg tablet</i> | 1 | |
| PHENOTHIAZINE DERIVATIVES | | |
| <i>promethazine hcl (12.5 mg supp.rect, 12.5 mg tablet, 25 mg supp.rect, 25 mg tablet, 50 mg tablet, 50 mg/ml ampul)</i> | 1 | |
| <i>promethazine hcl 50 mg/ml vial</i> | 1 | AL At least 2 yrs old |
| PROMETHEGAN (12.5 MG SUPPOS, 25 MG SUPPOSITORY) | 1 | |
| GASTROINTESTINAL DRUGS | | |
| ANTI-INFLAMMATORY AGENTS (GI DRUGS) | | |
| <i>alosetron hcl</i> | 3 | CC PA QPD 2.0 per day |
| <i>balsalazide disodium</i> | 1 | |
| <i>mesalamine (1.2 g tablet dr, 4 g/60 ml enema, 1000 mg supp.rect)</i> | 1 | |
| <i>mesalamine 800 mg tablet dr</i> | 3 | ST |
| ANTIDIARRHEA AGENTS | | |
| <i>diphenoxylate hcl/atropine 2.5-.025mg tablet</i> | 1 | |
| <i>loperamide hcl 2 mg capsule</i> | 1 | QL 30 / 30 days |
| VIBERZI | 3 | CC PA QPD 2.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|-------------------------|
| CATHARTICS AND LAXATIVES | | | |
| GAVILYTE-C | 1 | C \$0 | \$0 copay for age 45-75 |
| GAVILYTE-G | 1 | C \$0 | \$0 copay for age 45-75 |
| <i>peg 3350/sod sulf/sod bicarb/sod chloride/potassium chloride</i> | 1 | C \$0 | \$0 copay for age 45-75 |
| <i>peg 3350/sodium sulfate/sod chloride/kcl/ascorbate sod/vit c</i> | 1 | C \$0 | \$0 copay for age 45-75 |
| <i>sodium chloride/sodium bicarbonate/potassium chloride/peg</i> | 1 | C \$0 | \$0 copay for age 45-75 |
| <i>sodium sulfate/potassium sulfate/magnesium sulfate</i> | 1 | C \$0 | \$0 copay for age 45-75 |
| CHOLELITHOLYTIC AGENTS | | | |
| IQIRVO | 3 | CC PA QPD | 1.0 per day |
| LIVDELZI | 3 | CC PA QPD | 1.0 per day |
| OCALIVA | 3 | CC PA QPD | 1.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|-----------------------------|
| <i>ursodiol (250 mg tablet, 300 mg capsule, 500 mg tablet)</i> | 1 | | |
| DIGESTANTS | | | |
| CREON | 2 | | |
| ZENPEP | 2 | | |
| GI DRUGS, MISCELLANEOUS | | | |
| <i>dronabinol</i> | 3 | CC PA QPD | 3.0 per day |
| PROKINETIC AGENTS | | | |
| <i>metoclopramide hcl (5 mg tablet, 10 mg tablet)</i> | 1 | | |
| GENITOURINARY SMOOTH MUSCLE RELAXANTS | | | |
| ANTIMUSCARINICS | | | |
| <i>oxybutynin chloride (5 mg tab er 24, 5 mg tablet, 5 mg/5 ml syrup, 10 mg tab er 24, 15 mg tab er 24)</i> | 1 | | |
| <i>solifenacain succinate</i> | 3 | QL ST QPD | 90 / 90 days 1.0 per day |
| <i>tolterodine tartrate (2 mg cap er 24h, 4 mg cap er 24h)</i> | 3 | QL CC ST QPD | 30 / 30 days 1.0 per day |
| <i>tolterodine tartrate (1 mg tablet, 2 mg tablet)</i> | 3 | QL CC ST QPD | 60 / 30 days 2.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|-----------------------------|
| <i>trospium chloride 60 mg cap er 24h</i> | 3 | QL CC ST QPD | 30 / 30 days 1.0 per day |
| <i>trospium chloride 20 mg tablet</i> | 3 | QL CC ST QPD | 60 / 30 days 2.0 per day |

GOLD COMPOUNDS

| | | |
|--|---|-----------------------------|
| <i>auranofin</i> | 1 | |
| GONADOTROPINS AND ANTIGONADOTROPINS | | |
| ANTIGONADTROPINS | | |
| ORGOVYX | 3 | CC PA |
| ORIAHNN | 3 | CC PA QPD 2.0 per day |
| ORILISSA 150 MG TABLET | 3 | CC PA QPD 1.0 per day |
| ORILISSA 200 MG TABLET | 3 | CC PA QPD 2.0 per day |

GONADOTROPINS

| | | |
|----------------|---|----------|
| <i>ELIGARD</i> | 3 | CC PA |
|----------------|---|----------|

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| <i>leuprolide acetate (1 mg/0.2ml kit, 1 mg/0.2ml vial, 22.5 mg vial)</i> | 3 | CC | PA |
| LUPRON DEPOT (DEPOT-4 MONTH KIT, DEPOT 7.5 MG KIT, DEPOT 22.5 MG 3MO KIT, DEPOT 45 MG 6MO KIT) | 3 | CC | PA |
| LUPRON DEPOT 11.25 MG 3MO KIT | 3 | QL CC PA | 1 / 90 days |
| LUPRON DEPOT 3.75 MG KIT | 3 | QL CC PA | 1 / 30 days |
| HCV ANTIVIRALS | | | |
| HCV POLYMERASE INHIBITOR ANTIVIRALS | | | |
| <i>sofosbuvir/velpatasvir</i> | 3 | QL CC PA | 28 / 28 days |
| VOSEVI | 3 | QL CC PA | 28 / 28 days |
| HCV PROTEASE INHIBITOR ANTIVIRALS | | | |
| MAVYRET (50-20 MG PELLET PACKET, 100-40 MG TABLET) | 3 | QL CC PA | 84 / 28 days |
| HEAVY METAL ANTAGONISTS | | | |
| CHEMET | 3 | CC PA | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| <i>deferasirox (90 mg gran pack, 90 mg tablet, 125 mg tab disper, 180 mg gran pack, 180 mg tablet, 250 mg tab disper, 360 mg gran pack, 360 mg tablet, 500 mg tab disper)</i> | 3 | CC | PA |
| HORMONES AND SYNTHETIC SUBSTITUTES | | | |
| ADRENALS | | | |
| ARNUTITY ELLIPTA | 2 | QPD | 2.0 per day |
| ASMANEX | 2 | QPD | 0.07 per day |
| ASMANEX HFA | 2 | QPD | 0.9 per day |
| <i>budesonide (0.25mg/2ml ampul-neb, 0.5 mg/2ml ampul-neb)</i> | 1 | QPD | 4.0 per day |
| <i>budesonide 3 mg capdr - er</i> | 1 | | |
| <i>deflazacort (6 mg tablet, 18 mg tablet, 22.75mg/ml oral susp, 30 mg tablet, 36 mg tablet)</i> | 3 | CC | PA |
| <i>dexamethasone (0.5 mg tablet, 0.5 mg/5ml elixir, 0.5 mg/5ml solution, 0.75 mg tablet, 1 mg tablet, 1.5 mg tablet, 2 mg tablet, 4 mg tablet, 6 mg tablet)</i> | 1 | | |
| DEXAMETHASONE INTENSOL | 1 | | |
| <i>dexamethasone sodium phosphate 0.1 % drops</i> | 1 | | |
| <i>fludrocortisone acetate 0.1 mg tablet</i> | 1 | | |
| <i>fluticasone propionate (110 mcg aer w/adap, 220 mcg aer w/adap)</i> | 1 | QPD | 0.8 per day |
| <i>fluticasone propionate 44 mcg aer w/adap</i> | 1 | QPD | 0.71 per day |
| <i>fluticasone propionate (50 mcg blst w/dev, 100 mcg blst w/dev, 250 mcg blst w/dev)</i> | 1 | QPD | 4.0 per day |
| <i>hydrocortisone (5 mg tablet, 10 mg tablet, 20 mg tablet)</i> | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|---|
| ISTURISA 1 MG TABLET | 3 | <p>CC C Up to 60 mg per day allowed across all tablet strengths.</p> <p>PA QPD 8.0 per day</p> |
| ISTURISA 5 MG TABLET | 3 | <p>CC C Up to 60 mg per day allowed across all tablet strengths.</p> <p>PA QPD 12.0 per day</p> |
| MEDROL 2 MG TABLET | 2 | |
| <i>methylprednisolone (4 mg tab ds pk, 4 mg tablet, 8 mg tablet, 16 mg tablet, 32 mg tablet)</i> | 1 | |
| <i>prednisolone</i> | 1 | |
| <i>prednisone (1 mg tablet, 2.5 mg tablet, 5 mg tab ds pk, 5 mg tablet, 5 mg/5 ml solution, 10 mg tab ds pk, 10 mg tablet, 20 mg tablet, 50 mg tablet)</i> | 1 | |
| PREDNISONE INTENSOL | 1 | |
| PULMICORT FLEXHALER | 2 | QPD 0.07 per day |
| QVAR REDIHALER | 2 | QPD 0.8 per day |
| ANDROGENS | | |
| <i>danazol (50 mg capsule, 100 mg capsule, 200 mg capsule)</i> | 1 | |
| KYZATREX | 3 | <p>CC PA</p> |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|---------------|
| METHITEST | 2 | | |
| <i>testosterone (12.5/1.25g gel md pmp, 50 mg (1%) gel (gram))</i> | 3 | QL CC PA | 150 / 30 days |
| <i>testosterone (1.25g-1.62 gel packet, 2.5g-1.62% gel packet, 20.25/1.25 gel md pmp)</i> | 3 | QL CC PA | 75 / 30 days |
| <i>testosterone 25mg(1%) gel packet</i> | 3 | QL CC PA | 225 / 30 days |
| <i>testosterone 50 mg (1%) gel packet</i> | 3 | QL CC PA | 300 / 30 days |
| <i>testosterone cypionate (100 mg/ml vial, 200 mg/ml vial)</i> | 1 | QL | 5 / 30 days |
| <i>testosterone enanthate</i> | 1 | QL | 5 / 30 days |
| CONTRACEPTIVES | | | |
| AFIRMELLE | 1 | \$0 | |
| ALTAVERA | 1 | \$0 | |
| ALYACEN | 1 | \$0 | |
| AMETHIA | 1 | \$0 | |
| AMETHYST | 1 | \$0 | |
| ANNOVERA | 2 | \$0 | |
| APRI | 1 | \$0 | |
| ARANELLE | 1 | \$0 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---------------------|------|-----------------------|
| ASHLYNA | 1 | \$0 |
| AUBRA | 1 | \$0 |
| AUBRA EQ | 1 | \$0 |
| AUROVELA | 1 | \$0 |
| AUROVELA 24 FE | 1 | \$0 |
| AUROVELA FE | 1 | \$0 |
| AVERI | 2 | \$0 |
| AVIANE | 1 | \$0 |
| AYUNA | 1 | \$0 |
| AZURETTE | 1 | \$0 |
| BALZIVA | 1 | \$0 |
| BLISOVI 24 FE | 1 | \$0 |
| BLISOVI FE | 1 | \$0 |
| BRIELLYN | 1 | \$0 |
| CAMILA | 1 | \$0 |
| CAMRESE | 1 | \$0 |
| CAMRESE LO | 1 | \$0 |
| CAZIANT | 1 | \$0 |
| CHARLOTTE 24 FE | 1 | \$0 |
| CHATEAL EQ | 1 | \$0 |
| CRYSELLE | 1 | \$0 |
| CYRED | 1 | \$0 |
| CYRED EQ | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| DASETTA | 1 | \$0 |
| DAYSEE | 1 | \$0 |
| DEBLITANE | 1 | \$0 |
| <i>desogestrel-ethinyl estradiol/ethinyl estradiol</i> | 1 | \$0 |
| DOLISHALE | 1 | \$0 |
| <i>drosipренон/этиныл эстрадиол/левомекофолат кальция</i> | 1 | \$0 |
| ELINEST | 1 | \$0 |
| ELLA | 2 | \$0 |
| ELURYNG | 1 | \$0 |
| EMZAHH | 1 | \$0 |
| ENILLORING | 1 | \$0 |
| ENPRESSE | 1 | \$0 |
| ENSKYCE | 1 | \$0 |
| ERRIN | 1 | \$0 |
| ESTARYLLA | 1 | \$0 |
| <i>ethинил эстрадиол/дросипренон</i> | 1 | \$0 |
| <i>этинонодиол дикарбонат-этиныл эстрадиол</i> | 1 | \$0 |
| <i>етоногестрел/этиныл эстрадиол</i> | 1 | \$0 |
| FALMINA | 1 | \$0 |
| FEIRZA | 1 | \$0 |
| FEMLYV | 1 | \$0 |
| FINZALA | 1 | \$0 |
| GALBRIELA | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---------------------|------|-----------------------|
| GEMMILY | 1 | \$0 |
| HAILEY | 1 | \$0 |
| HAILEY 24 FE | 1 | \$0 |
| HAILEY FE | 1 | \$0 |
| HALOETTE | 1 | \$0 |
| HEATHER | 1 | \$0 |
| ICLEVIA | 1 | \$0 |
| INCASSIA | 1 | \$0 |
| ISIBLOOM | 1 | \$0 |
| JAIMIESS | 1 | \$0 |
| JASMIEL | 1 | \$0 |
| JENCYCLA | 1 | \$0 |
| JOLESSA | 1 | \$0 |
| JOYEAUX | 1 | \$0 |
| JULEBER | 1 | \$0 |
| JUNEL | 1 | \$0 |
| JUNEL FE | 1 | \$0 |
| JUNEL FE 24 | 1 | \$0 |
| KAITLIB FE | 1 | \$0 |
| KALLIGA | 1 | \$0 |
| KARIVA | 1 | \$0 |
| KELNOR 1-35 | 1 | \$0 |
| KELNOR 1-50 | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| KURVELO | 1 | \$0 |
| LARIN | 1 | \$0 |
| LARIN 24 FE | 1 | \$0 |
| LARIN FE | 1 | \$0 |
| LAYOLIS FE | 1 | \$0 |
| LEENA | 1 | \$0 |
| LESSINA | 1 | \$0 |
| LEVONEST | 1 | \$0 |
| <i>levonorgestrel/ethinyl estradiol (levonorgestrel/ethin.estradiol 0.1-0.02mg tablet, levonorgestrel/ethin.estradiol 0.15-0.03 tablet, levonorgestrel/ethin.estradiol 0.15-0.03 tbdspk 3mo, levonorgestrel/ethin.estradiol 6-5-10 tablet, levonorgestrel/ethin.estradiol 90-20 mcg tablet)</i> | 1 | \$0 |
| <i>levonorgestrel/ethinyl estradiol and ethinyl estradiol</i> | 1 | \$0 |
| <i>levonorgestrel/ethinyl estradiol/iron</i> | 1 | \$0 |
| LEVORA-28 | 1 | \$0 |
| LO LOESTRIN FE | 2 | \$0 |
| LO-ZUMANDIMINE | 1 | \$0 |
| LOESTRIN | 1 | \$0 |
| LOESTRIN FE 1.5-30 TABLET | 1 | \$0 |
| LOJAIMIESS | 1 | \$0 |
| LORYNA | 1 | \$0 |
| LOW-OGESTREL | 1 | \$0 |
| LUTERA | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| LYLEQ | 1 | \$0 |
| LYZA | 1 | \$0 |
| MARLISSA | 1 | \$0 |
| MELEYA | 1 | \$0 |
| MERZEE | 1 | \$0 |
| MIBELAS 24 FE | 1 | \$0 |
| MICROGESTIN | 1 | \$0 |
| MICROGESTIN FE | 1 | \$0 |
| MILI | 1 | \$0 |
| MINZOYA | 1 | \$0 |
| MONO-LINYAH | 1 | \$0 |
| NATAZIA | 2 | \$0 |
| NECON | 1 | \$0 |
| NEXTSTELLIS | 2 | \$0 |
| NIKKI | 1 | \$0 |
| NORA-BE | 1 | \$0 |
| <i>norelgestromin/ethynodiolide</i> | 1 | \$0 |
| <i>norethindrone 0.35 mg tablet</i> | 1 | \$0 |
| <i>norethindrone acetate-ethynodiolide (1mg-20mcg tablet, 1.5-0.03mg tablet)</i> | 1 | \$0 |
| <i>norethindrone acetate-ethynodiolide/ferrous fumarate (1mg-20(21) tablet, 1mg-20(24) capsule, 1mg-20(24) tab chew, 1.5-30(21) tablet, 5-7-9-7 tablet)</i> | 1 | \$0 |
| <i>norethindrone-ethynodiolide/ferrous fumarate</i> | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| <i>norgestimate-ethynodiol (0.25-0.035 tablet, 7daysx3 28 tablet, 7daysx3 lo tablet)</i> | 1 | \$0 |
| NORTREL | 1 | \$0 |
| NYLIA | 1 | \$0 |
| NYMYO | 1 | \$0 |
| OCELLA | 1 | \$0 |
| ORQUIDEA | 1 | \$0 |
| PHILITH | 1 | \$0 |
| PIMTREA | 1 | \$0 |
| PORTIA | 1 | \$0 |
| RECLIPSEN | 1 | \$0 |
| RIVELSA | 1 | \$0 |
| ROSYRAH | 1 | \$0 |
| SETLAKIN | 1 | \$0 |
| SHAROBEL | 1 | \$0 |
| SIMLIYA | 1 | \$0 |
| SIMPESSE | 1 | \$0 |
| SLYND | 2 | \$0 |
| SPRINTEC | 1 | \$0 |
| SRONYX | 1 | \$0 |
| SYEDA | 1 | \$0 |
| TARINA 24 FE | 1 | \$0 |
| TARINA FE | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---------------------|------|-----------------------|
| TARINA FE 1-20 EQ | 1 | \$0 |
| TAYSOFY | 1 | \$0 |
| TILIA FE | 1 | \$0 |
| TRI-ESTARYLLA | 1 | \$0 |
| TRI-LEGEST FE | 1 | \$0 |
| TRI-LINYAH | 1 | \$0 |
| TRI-LO-ESTARYLLA | 1 | \$0 |
| TRI-LO-MARZIA | 1 | \$0 |
| TRI-LO-MILI | 1 | \$0 |
| TRI-LO-SPRINTEC | 1 | \$0 |
| TRI-MILI | 1 | \$0 |
| TRI-NYMYO | 1 | \$0 |
| TRI-SPRINTEC | 1 | \$0 |
| TRI-VYLIBRA | 1 | \$0 |
| TRI-VYLIBRA LO | 1 | \$0 |
| TRIVORA-28 | 1 | \$0 |
| TULANA | 1 | \$0 |
| TUROQZ | 1 | \$0 |
| TWIRLA | 2 | \$0 |
| TYDEMY | 1 | \$0 |
| VALTYA | 1 | \$0 |
| VELIVET | 2 | \$0 |
| VESTURA | 1 | \$0 |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| VIENVA | 1 | \$0 |
| VIORELE | 1 | \$0 |
| VOLNEA | 1 | \$0 |
| VYFEMLA | 1 | \$0 |
| VYLIBRA | 1 | \$0 |
| WERA | 1 | \$0 |
| WYMZYA FE | 1 | \$0 |
| XARAH FE | 1 | \$0 |
| XELRIA FE | 1 | \$0 |
| XULANE | 1 | \$0 |
| ZAFEMY | 1 | \$0 |
| ZARAH | 1 | \$0 |
| ZOVIA 1-35 | 1 | \$0 |
| ZUMANDIMINE | 1 | \$0 |
| PITUITARY | | |
| <i>desmopressin acetate 10/spray spray/pump</i> | 1 | QPD 0.5 per day |
| <i>desmopressin acetate (0.1 mg tablet, 0.2 mg tablet)</i> | 1 | |
| <i>desmopressin acetate (non-refrigerated)</i> | 1 | QPD 0.5 per day |
| PROGESTINS | | |
| CRINONE 8% GEL | 3 | CC PA |
| DEPO-SUBQ PROVERA 104 | 2 | \$0 |
| ENDOMETRIN | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| GALLIFREY | 1 | | |
| <i>medroxyprogesterone acetate (2.5 mg tablet, 5 mg tablet, 10 mg tablet)</i> | 1 | | |
| <i>medroxyprogesterone acetate (150 mg/ml syringe, 150 mg/ml vial)</i> | 1 | \$0 | |
| <i>megestrol acetate (20 mg tablet, 40 mg tablet, 400mg/10ml oral susp)</i> | 1 | | |
| <i>norethindrone acetate 5 mg tablet</i> | 1 | | |
| <i>progesterone, micronized (100 mg capsule, 200 mg capsule)</i> | 1 | QL | 30 / 30 days |
| IMMUNOMODULATORY AGENTS (90:00) | | | |
| COMPLEMENT INHIBITOR AGENTS (90:20) | | | |
| TAVNEOS | 3 | CC PA QPD | 6.0 per day |
| INSULINS | | | |
| LONG-ACTING INSULINS | | | |
| <i>insulin degludec (100/ml (3) insulin pen, 100/ml vial, 200/ml (3) insulin pen)</i> | 3 | CC ST | |
| <i>insulin glargine, human recombinant analog (100/ml (3) insulin pen, 100/ml vial)</i> | 1 | | |
| <i>insulin glargine-yfgn (100/ml (3) insulin pen, 100/ml vial)</i> | 1 | | |
| REZVOGLAR KWIKPEN | 1 | | |
| RAPID-ACTING INSULINS | | | |
| ADMELOG | 2 | | |
| ADMELOG SOLOSTAR | 2 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| <i>insulin aspart protamine human/insulin aspart (art prot/insulin 70-30/ml insulin pen, art prot/insulin 70-30/ml vial)</i> | 1 | | |
| <i>insulin lispro (100/ml insulin pen, 100/ml vial)</i> | 1 | | |
| <i>insulin lispro protamine and insulin lispro</i> | 1 | | |
| SHORT-ACTING INSULINS | | | |
| HUMULIN R U-500 | 3 | ST | |
| HUMULIN R U-500 KWIKPEN | 3 | ST | |
| INTERLEUKIN-MEDIATED AGENTS | | | |
| INTERLEUKIN-MEDIATED AGENTS, MISC | | | |
| ACTEMRA 162 MG/0.9 ML SYRINGE | 3 | CC PA QPD | 0.13 per day |
| ACTEMRA ACTPEN | 3 | CC PA QPD | 0.13 per day |
| COSENTYX (2 SYRINGES) | 3 | CC PA QPD | 0.08 per day |
| COSENTYX SENSOREADY (2 PENS) | 3 | CC PA QPD | 0.08 per day |
| COSENTYX SENSOREADY PEN | 3 | CC PA QPD | 0.08 per day |
| COSENTYX SYRINGE | 3 | CC PA QPD | 0.08 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|--------------|-------------------|
| COSENTYX UNOREADY PEN | 3 | CC | PA | QPD 0.08 per day |
| STELARA (45 MG/0.5 ML SYRINGE, 45 MG/0.5 ML VIAL, 90 MG/ML SYRINGE) | 3 | CC | PA | QPD 0.018 per day |
| ION-REMOVING AGENTS | | | | |
| PHOSPHATE-REMOVING AGENTS | | | | |
| <i>calcium acetate</i> | 1 | | | |
| <i>lanthanum carbonate</i> | 3 | CC | PA | QPD 3.0 per day |
| <i>sevelamer carbonate 0.8 g powd pack</i> | 3 | CC | PA | QPD 6.0 per day |
| <i>sevelamer carbonate 2.4 g powd pack</i> | 3 | CC | PA | QPD 3.0 per day |
| <i>sevelamer carbonate 800 mg tablet</i> | 1 | | | |
| <i>sevelamer hcl</i> | 3 | CC | PA | QPD 9.0 per day |
| POTASSIUM-REMOVING AGENTS | | | | |
| LOKELMA | 2 | QL | 35 / 30 days | |
| <i>sodium polystyrene sulfonate</i> | 1 | | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|------------------------------|--|
| SPS 15 GM/60 ML SUSPENSION | 2 | | |
| VELTASSA (8.4 GM POWDER PACKET, 16.8 GM POWDER PACKET, 25.2 GM POWDER PACKET) | 3 | ST | |
| JANUS KINASE INHIBITORS (90:24) | | | |
| JANUS KINASE INHIBITORS, MISCELLANEOUS | | | |
| OLUMIANT | 3 | CC PA QPD 1.0 per day | |
| XELJANZ (5 MG TABLET, 10 MG TABLET) | 3 | CC PA QPD 2.0 per day | |
| XELJANZ XR | 3 | CC PA QPD 1.0 per day | |
| KALLIKREIN-KININ SYSTEM INHIBITORS | | | |
| KALLIKREIN | | | |
| ORLADEYO | 3 | CC PA QPD 1.0 per day | |
| TAKHYRO 300 MG/2 ML VIAL | 3 | CC PA QPD 0.15 per day | |
| MACROLIDE ANTIBIOTICS | | | |
| ERYTHROMYCIN ANTIBIOTICS | | | |
| ERYTHROCIN STEARATE | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------------|
| <i>erythromycin base (250 mg capsule dr, 250 mg tablet, 250 mg tablet dr, 333 mg tablet dr, 500 mg tablet, 500 mg tablet dr)</i> | 1 | |
| <i>erythromycin ethylsuccinate 400 mg tablet</i> | 1 | |
| OTHER MACROLIDE ANTIBIOTICS | | |
| <i>azithromycin (1 g packet, 250 mg tablet, 500 mg tablet, 600 mg tablet)</i> | 1 | |
| <i>clarithromycin (250 mg tablet, 500 mg tablet)</i> | 1 | |
| <i>fidaxomicin</i> | 3 | CC PA |
| MINERALOCORTICOID (ALDOSTERONE) ANTAGENTS | | |
| STEROIDAL MINERALOCORTICOID RECEPTOR ANT | | |
| <i>eplerenone</i> | 3 | ST |
| <i>spironolactone (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| <i>spironolactone/hydrochlorothiazide</i> | 1 | |
| MISC. BETA-LACTAM ANTIBIOTICS | | |
| MONOBACTAM ANTIBIOTICS | | |
| <i>CAYSTON</i> | 3 | CC PA QPD 1.5 per day |
| MISCELLANEOUS THERAPEUTIC AGENTS | | |
| 5-ALPHA-REDUCTASE INHIBITORS (92:04) | | |
| <i>dutasteride 0.5 mg capsule</i> | 1 | |
| <i>finasteride 5 mg tablet</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|-------------|--|
| ANTIGOUT AGENTS | | | | |
| <i>allopurinol (100 mg tablet, 300 mg tablet)</i> | 1 | | | |
| <i>febuxostat</i> | 3 | CC PA QPD | 1.0 per day | |
| BONE RESORPTION INHIBITORS | | | | |
| <i>alendronate sodium (5 mg tablet, 10 mg tablet, 35 mg tablet, 70 mg tablet)</i> | 1 | | | |
| <i>ibandronate sodium 150 mg tablet</i> | 1 | | | |
| OTHER MISCELLANEOUS THERAPEUTIC AGENTS | | | | |
| <i>betaine</i> | 1 | | | |
| <i>EVRYSDI 60 MG/80 ML(0.75MG/ML)</i> | 3 | CC PA | | |
| <i>glutamine 5 g powd pack</i> | 3 | CC PA | | |
| <i>levocarnitine 100 mg/ml solution</i> | 1 | | | |
| <i>levocarnitine (with sugar) 100 mg/ml solution</i> | 1 | | | |
| <i>REZUROCK</i> | 3 | CC PA | | |
| <i>SKYCLARYS</i> | 3 | CC PA QPD | 3.0 per day | |
| PROTECTIVE AGENTS | | | | |
| <i>dalfampridine 10 mg tab er 12h</i> | 3 | CC PA QPD | 2.0 per day | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|------------------------------|
| MTOR INHIBITORS | | |
| MTOR INHIBITORS, MISCELLANEOUS | | |
| <i>sirolimus (0.5 mg tablet, 1 mg tablet, 2 mg tablet)</i> | 1 | |
| MULTIPLE SCLEROSIS AGENTS | | |
| AMINO ACID POLYMERS | | |
| <i>glatiramer acetate 20 mg/ml syringe</i> | 3 | CC PA QPD 1.0 per day |
| <i>glatiramer acetate 40 mg/ml syringe</i> | 3 | CC PA QPD 0.43 per day |
| GLATOPA 20 MG/ML SYRINGE | 3 | CC PA QPD 1.0 per day |
| GLATOPA 40 MG/ML SYRINGE | 3 | CC PA QPD 0.43 per day |
| ANTIMETABOLITES | | |
| MAVENCLAD | 3 | CC PA |
| FUMARATES | | |
| <i>dimethyl fumarate (120 mg capsule dr, 240 mg capsule dr)</i> | 3 | CC PA QPD 2.0 per day |
| <i>dimethyl fumarate 120-240 mg capsule dr</i> | 3 | CC PA |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| INTERFERONS | | | |
| AVONEX (4 PACK) | 3 | CC PA QPD | 0.08 per day |
| AVONEX PEN (4 PACK) | 3 | CC PA QPD | 0.08 per day |
| BETASERON (0.3 MG KIT, 0.3 MG VIAL) | 3 | CC PA QPD | 0.5 per day |
| EXTAVIA (0.3 MG KIT, 0.3 MG VIAL) | 3 | CC PA QPD | 0.5 per day |
| REBIF | 3 | CC PA QPD | 0.22 per day |
| REBIF REBIDOSE | 3 | CC PA | |
| SPHINGOSINE 1-PHOSPHATE (S1P) AGENTS | | | |
| <i>fingolimod hcl</i> | 3 | CC PA QPD | 1.0 per day |
| GILENYA 0.25 MG CAPSULE | 3 | CC PA QPD | 1.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| NONHORMONAL CONTRACEPTIVES | | |
| <i>diaphragms</i> | 1 | \$0 |
| PHEXXI | 1 | \$0 |
| NONSTEROIDAL ANTI-INFLAMMATORY AGENTS | | |
| CYCLOOXYGENASE-2 (COX-2) INHIBITORS | | |
| <i>celecoxib (50 mg capsule, 100 mg capsule, 200 mg capsule, 400 mg capsule)</i> | 1 | |
| REVERSIBLE COX-1/COX-2 INHIBITORS | | |
| <i>diclofenac potassium 50 mg tablet</i> | 1 | |
| <i>diclofenac sodium 1 % gel (gram)</i> | 1 | QL 300 / 30 days |
| <i>diclofenac sodium (25 mg tablet dr, 50 mg tablet dr, 75 mg tablet dr, 100 mg tab er 24h)</i> | 1 | |
| <i>diclofenac sodium/misoprostol</i> | 1 | |
| <i>etodolac (200 mg capsule, 300 mg capsule, 400 mg tab er 24h, 400 mg tablet, 500 mg tab er 24h, 500 mg tablet, 600 mg tab er 24h)</i> | 1 | |
| <i>flurbiprofen 100 mg tablet</i> | 1 | |
| IBU | 1 | |
| <i>ibuprofen (100 mg/5ml oral susp, 400 mg tablet, 600 mg tablet, 800 mg tablet)</i> | 1 | |
| <i>indomethacin (25 mg capsule, 50 mg capsule, 50 mg supp.rect, 75 mg capsule er)</i> | 1 | |
| <i>ketorolac tromethamine 10 mg tablet</i> | 1 | QL 20 / 30 days |
| <i>meloxicam (7.5 mg tablet, 15 mg tablet)</i> | 1 | |
| <i>nabumetone (500 mg tablet, 750 mg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | | | |
|---|------|-----------------------|---------------|--|--|--|
| <i>naproxen (250 mg tablet, 375 mg tablet, 375 mg tablet dr, 500 mg tablet, 500 mg tablet dr)</i> | 1 | | | | | |
| <i>naproxen sodium (275 mg tablet, 550 mg tablet)</i> | 1 | | | | | |
| <i>oxaprozin</i> | 1 | | | | | |
| <i>piroxicam (10 mg capsule, 20 mg capsule)</i> | 1 | | | | | |
| <i>sulindac (150 mg tablet, 200 mg tablet)</i> | 1 | | | | | |
| SALICYLATES | | | | | | |
| <i>aspirin/dipyridamole</i> | 1 | QL | 60 / 30 days | | | |
| <i>butalbital/aspirin/caffeine 50-325-40 tablet</i> | 3 | CC PA QPD | 2.0 per day | | | |
| <i>salsalate (500 mg tablet, 750 mg tablet)</i> | 1 | | | | | |
| OXYTOCICS | | | | | | |
| <i>methylergonovine maleate 0.2 mg tablet</i> | 1 | QL | 30 / 30 days | | | |
| <i>mifepristone 200 mg tablet</i> | 1 | HCR \$0 | | | | |
| PARATHYROID AND ANTIPARATHYROID AGENTS | | | | | | |
| ANTIPARATHYROID AGENTS | | | | | | |
| <i>calcitonin, salmon, synthetic 200/spray spray/pump</i> | 1 | | | | | |
| <i>cinacalcet hcl</i> | 1 | CC QPD | 4.0 per day | | | |
| PARATHYROID AGENTS | | | | | | |
| <i>teriparatide</i> | 3 | CC PA QPD | 0.083 per day | | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|--|------|-----------------------|----|-------------------|
| TYMLOS | 3 | CC | PA | QPD 0.052 per day |
| PENICILLIN ANTIBIOTICS | | | | |
| AMINOPENICILLIN ANTIBIOTICS | | | | |
| <i>amoxicillin (250 mg capsule, 500 mg capsule, 500 mg tablet, 875 mg tablet)</i> | 1 | | | |
| <i>amoxicillin/potassium clavulanate (amoxicillin/potassium 250-125 mg tablet, amoxicillin/potassium 500-125 mg tablet, amoxicillin/potassium 875-125 mg tablet, amoxicillin/potassium 1000-62.5 tab er 12h)</i> | 1 | | | |
| <i>ampicillin trihydrate</i> | 1 | | | |
| NATURAL PENICILLIN ANTIBIOTICS | | | | |
| <i>penicillin v potassium (250 mg tablet, 500 mg tablet)</i> | 1 | | | |
| PENICILLINASE-RESISTANT PENICILLINS | | | | |
| <i>dicloxacillin sodium</i> | 1 | | | |
| PHOSPHODIESTERASE-4 INHIBITORS (90:24) | | | | |
| PHOSPHODIESTERASE-4 INHIBITORS, MISC | | | | |
| OTEZLA (10-20-30MG START 14 DAY, 10-20-30MG START 28 DAY, 30 MG TABLET) | 3 | CC | PA | QPD 2.0 per day |
| OTEZLA (10-20 MG STARTER 28 DAY, 20 MG TABLET) | 3 | CC | PA | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|-----------------------|
| RENIN-ANGIOTENSIN-ALDOSTERONE SYS. INHIB ANGIOTENSIN II RECEPTOR ANTAGONIST/NEPROLYS | | |
| <i>sacubitril/valsartan</i> | 1 | |
| ANGIOTENSIN II RECEPTOR ANTAGONISTS | | |
| <i>irbesartan</i> | 1 | |
| <i>irbesartan/hydrochlorothiazide</i> | 1 | |
| <i>losartan potassium (25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| <i>losartan potassium/hydrochlorothiazide</i> | 1 | |
| <i>olmesartan medoxomil</i> | 1 | |
| <i>telmisartan</i> | 1 | |
| <i>valsartan (40 mg tablet, 80 mg tablet, 160 mg tablet, 320 mg tablet)</i> | 1 | |
| <i>valsartan/hydrochlorothiazide</i> | 1 | |
| ANGIOTENSIN-CONVERTING ENZYME INHIBITORS | | |
| <i>benazepril hcl (5 mg tablet, 10 mg tablet, 20 mg tablet, 40 mg tablet)</i> | 1 | |
| <i>benazepril hcl/hydrochlorothiazide</i> | 1 | |
| <i>captopril (12.5 mg tablet, 25 mg tablet, 50 mg tablet, 100 mg tablet)</i> | 1 | |
| <i>enalapril maleate (2.5 mg tablet, 5 mg tablet, 10 mg tablet, 20 mg tablet)</i> | 1 | |
| <i>enalapril maleate/hydrochlorothiazide</i> | 1 | |
| <i>lisinopril (2.5 mg tablet, 5 mg tablet, 10 mg tablet, 20 mg tablet, 30 mg tablet, 40 mg tablet)</i> | 1 | |
| <i>lisinopril/hydrochlorothiazide</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|----|-----------------|
| MINERALOCORTICOID (ALDOSTERONE) ANTAGNTS | | | | |
| KERENDIA (10 MG TABLET, 20 MG TABLET) | 3 | CC | PA | QPD 1.0 per day |
| RESPIRATORY TRACT AGENTS | | | | |
| ANTIFIBROTIC AGENTS | | | | |
| <i>pirfenidone 267 mg tablet</i> | 3 | CC | PA | QPD 6.0 per day |
| <i>pirfenidone 534 mg tablet</i> | 3 | CC | PA | |
| <i>pirfenidone 801 mg tablet</i> | 3 | CC | PA | QPD 3.0 per day |
| ANTITUSSIVES | | | | |
| <i>benzonatate</i> | 1 | | | |
| <i>promethazine hcl/codeine 6.25-10/5 syrup</i> | 1 | | | |
| <i>promethazine hcl/dextromethorphan hbr</i> | 1 | | | |
| MUCOLYTIC AGENTS | | | | |
| PULMOZYME | 3 | CC | PA | QPD 2.5 per day |
| PHOSPHODIESTERASE TYPE 4 INHIBITORS | | | | |
| <i>roflumilast</i> | 3 | CC | PA | QPD 1.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|--|------|-----------------------|------------------------------|--|
| VASODILATING AGENTS (RESPIRATORY TRACT) | | | | |
| ADEMPAS | 3 | QL CC PA | 90 / 30 days | |
| <i>ambrisentan</i> | 3 | QL CC PA | 30 / 30 days | |
| <i>treprostinil sodium</i> | 3 | CC PA | | |
| TYVASO | 3 | CC PA QPD | 2.9 per day | |
| TYVASO INSTITUTIONAL START KIT | 3 | QL CC PA | 81.2 / 28 days | |
| TYVASO REFILL KIT | 3 | CC PA QPD | 2.9 per day | |
| TYVASO STARTER KIT | 3 | QL CC PA | 81.2 / 28 days | |
| UPTRAVI 200-800 TITRATION PACK | 3 | QL CC PA QPD | 200 / 28 days 2.0 per day | |
| UPTRAVI (400 MCG TABLET, 600 MCG TABLET, 800 MCG TABLET, 1,000 MCG TABLET, 1,200 MCG TABLET, 1,400 MCG TABLET, 1,600 MCG TABLET) | 3 | CC PA QPD | 2.0 per day | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|------------------------------|
| UPTRAVI 200 MCG TABLET | 3 | QL CC PA QPD | 140 / 28 days 2.0 per day |
| SKELETAL MUSCLE RELAXANTS | | | |
| CENTRALLY ACTING SKELETAL MUSCLE RELAXNT | | | |
| cyclobenzaprine hcl (5 mg tablet, 10 mg tablet) | 1 | QL | 90 / 30 days |
| methocarbamol (500 mg tablet, 750 mg tablet) | 1 | QL | 120 / 30 days |
| tizanidine hcl (2 mg tablet, 4 mg tablet) | 1 | QL | 90 / 30 days |
| GABA-DERIVATIVE SKELETAL MUSCLE RELAXANT | | | |
| baclofen (5 mg tablet, 10 mg tablet, 20 mg tablet) | 1 | | |
| SKIN AND MUCOUS MEMBRANE AGENTS | | | |
| ANTIPROLIFERANTS | | | |
| fluorouracil 5 % cream (g) | 1 | QL | 1 / 365 days |
| imiquimod 5 % cream pack | 1 | AL | At least 12 yrs old |
| ANTIPRURITICS AND LOCAL ANESTHETICS | | | |
| lidocaine 5 % adh. patch | 1 | | |
| lidocaine 5 % oint. (g) | 1 | QL | 60 / 30 days |
| lidocaine/prilocaine 2.5 %-2.5% cream (g) | 1 | | |
| phenazopyridine hcl (100 mg tablet, 200 mg tablet) | 1 | | |
| ASTRINGENTS (84:12) | | | |
| DRYSOL | 2 | QL | 75 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|--|------|-----------------------|------------------|--|
| CELL STIMULANTS AND PROLIFERANTS | | | | |
| <i>tretinoin (0.025 % cream (g), 0.05 % cream (g), 0.1 % cream (g))</i> | 1 | QL | 20 / 30 days | |
| | | CC | | |
| | | AL | Up to 30 yrs old | |
| | | QPD | 0.7 per day | |
| <i>tretinoin (0.01 % gel (gram), 0.025 % gel (gram))</i> | 1 | QL | 15 / 30 days | |
| | | CC | | |
| | | AL | Up to 30 yrs old | |
| | | QPD | 0.5 per day | |
| KERATOLYTIC AGENTS | | | | |
| ACCUTANE | 3 | QL | 60 / 30 days | |
| | | CC | | |
| | | PA | | |
| <i>acitretin</i> | 3 | CC | | |
| | | PA | | |
| | | QPD | 2.0 per day | |
| <i>adapalene 0.3 % gel (gram)</i> | 1 | CC | | |
| | | AL | Up to 30 yrs old | |
| | | QPD | 1.5 per day | |
| AMNESTEEM | 3 | QL | 60 / 30 days | |
| | | CC | | |
| | | PA | | |
| CLARAVIS | 3 | QL | 60 / 30 days | |
| | | CC | | |
| | | PA | | |
| <i>isotretinoin (10 mg capsule, 20 mg capsule, 25 mg capsule, 30 mg capsule, 35 mg capsule, 40 mg capsule)</i> | 3 | QL | 60 / 30 days | |
| | | CC | | |
| | | PA | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|--|------|-----------------------|--------------|
| <i>podofilox 0.5 % solution</i> | 1 | | |
| ZENATANE | 3 | QL CC PA | 60 / 30 days |
| SKIN AND MUCOUS MEMBRANE AGENTS, MISC. | | | |
| DUPIXENT 200 MG/1.14 ML PEN | 3 | CC PA QPD | 0.09 per day |
| DUPIXENT 300 MG/2 ML PEN | 3 | CC PA QPD | 0.15 per day |
| DUPIXENT 100 MG/0.67 ML SYRING | 3 | CC PA QPD | 0.05 per day |
| DUPIXENT 200 MG/1.14 ML SYRING | 3 | CC PA QPD | 0.09 per day |
| DUPIXENT 300 MG/2 ML SYRINGE | 3 | CC PA QPD | 0.15 per day |
| SMOOTH MUSCLE RELAXANTS | | | |
| RESPIRATORY SMOOTH MUSCLE RELAXANTS | | | |
| <i>theophylline anhydrous (300 mg tab er 12h, 400 mg tab er 24h, 450 mg tab er 12h, 600 mg tab er 24h)</i> | 1 | | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|----|------------------|
| SOMATOSTATIN AGONISTS AND ANTAGONISTS | | | | |
| SOMATOSTATIN AGONISTS | | | | |
| MYCAPSSA | 3 | CC | PA | QPD 4.0 per day |
| <i>octreotide acetate (100 mcg/ml ampul, 100 mcg/ml vial)</i> | 3 | CC | PA | QPD 15.0 per day |
| <i>octreotide acetate (500 mcg/ml ampul, 500 mcg/ml vial)</i> | 3 | CC | PA | QPD 3.0 per day |
| <i>octreotide acetate (50 mcg/ml syringe, 100 mcg/ml syringe, 500 mcg/ml syringe)</i> | 3 | CC | PA | |
| <i>octreotide acetate (50 mcg/ml ampul, 50 mcg/ml vial)</i> | 3 | CC | PA | QPD 30.0 per day |
| <i>octreotide acetate 1000mcg/ml vial</i> | 3 | CC | PA | QPD 1.5 per day |
| <i>octreotide acetate 200 mcg/ml vial</i> | 3 | CC | PA | QPD 7.5 per day |
| SOMATOTROPIN AGONISTS AND ANTAGONISTS | | | | |
| SOMATOTROPIN ANTAGONISTS | | | | |
| SOMAVERT | 3 | CC | PA | QPD 1.0 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| SYMPATHOMIMETIC (ADRENERGIC) AGENTS ALPHA- AND BETA-ADRENERGIC AGONISTS | | |
| <i>epinephrine (0.15/0.15 auto inject, 0.15mg/0.3 auto inject, 0.3mg/0.3 auto inject)</i> | 1 | QL 6 / 365 days |
| ALPHA-ADRENERGIC AGONISTS | | |
| <i>midodrine hcl</i> | 1 | |
| THYROID AND ANTITHYROID AGENTS | | |
| ANTITHYROID AGENTS | | |
| <i>methimazole (5 mg tablet, 10 mg tablet)</i> | 1 | |
| <i>propylthiouracil 50 mg tablet</i> | 1 | |
| THYROID AGENTS | | |
| ADTHYZA (15 MG TABLET, 30 MG TABLET, 60 MG TABLET, 90 MG TABLET, 120 MG TABLET) | 2 | |
| ARMOUR THYROID | 2 | |
| EUTHYROX | 2 | |
| LEVO-T | 2 | |
| <i>levothyroxine sodium (13 mcg capsule, 25 mcg capsule, 25 mcg tablet, 50 mcg capsule, 50 mcg tablet, 75 mcg capsule, 75 mcg tablet, 88 mcg capsule, 88 mcg tablet, 100 mcg capsule, 100 mcg tablet, 112 mcg capsule, 112 mcg tablet, 125 mcg capsule, 125 mcg tablet, 137 mcg capsule, 137 mcg tablet, 150 mcg capsule, 150 mcg tablet, 175 mcg capsule, 175 mcg tablet, 200 mcg capsule, 200 mcg tablet, 300 mcg tablet)</i> | 1 | |
| LEVOXYL | 2 | |
| <i>liothyronine sodium (5 mcg tablet, 25 mcg tablet, 50 mcg tablet)</i> | 1 | |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|--|------|------------------------------|
| NIVA THYROID | 2 | |
| NP THYROID | 2 | |
| RENTHYROID | 2 | |
| SYNTHROID | 2 | |
| <i>thyroid, pork (15 mg tablet, 30 mg tablet, 60 mg tablet, 90 mg tablet, 120 mg tablet)</i> | 1 | |
| TIROSINT (13 MCG CAPSULE, 25 MCG CAPSULE, 50 MCG CAPSULE, 75 MCG CAPSULE, 88 MCG CAPSULE, 100 MCG CAPSULE, 112 MCG CAPSULE, 125 MCG CAPSULE, 137 MCG CAPSULE, 150 MCG CAPSULE, 175 MCG CAPSULE, 200 MCG CAPSULE) | 2 | |
| TIROSINT (37.5 MCG CAPSULE, 44 MCG CAPSULE, 62.5 MCG CAPSULE) | 2 | |
| TIROSINT-SOL | 2 | |
| UNITHROID | 2 | |
| TUMOR NECROSIS FACTOR INHIBITORS | | |
| TUMOR NECROSIS FACTOR INHIBITORS, MISC | | |
| <i>adalimumab-aaty (20mg/0.2ml syringe/kit, 40mg/0.4ml autoinj/kit, 40mg/0.4ml syringe/kit, 80mg/0.8ml autoinj/kit)</i> | 3 | CC PA QPD 0.15 per day |
| <i>adalimumab-adaz (10mg/0.1ml syringe, 20mg/0.2ml syringe, 40mg/0.4ml pen injctr, 40mg/0.4ml syringe, 80mg/0.8ml pen injctr)</i> | 3 | CC PA QPD 0.15 per day |
| ENBREL (25 MG/0.5 ML SYRINGE, 25 MG/0.5 ML VIAL, 50 MG/ML SYRINGE) | 3 | CC PA QPD 0.15 per day |
| ENBREL MINI | 3 | CC PA QPD 0.15 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|--|------|-----------------------|----|------------------|
| ENBREL SURECLICK | 3 | CC | PA | QPD 0.15 per day |
| HADLIMA | 3 | CC | PA | QPD 0.15 per day |
| HADLIMA PUSHTOUCH | 3 | CC | PA | QPD 0.15 per day |
| HADLIMA(CF) | 3 | CC | PA | QPD 0.15 per day |
| HADLIMA(CF) PUSHTOUCH | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA PEN | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA(CF) (10 MG/0.1 ML SYRING, 20 MG/0.2 ML SYRING, 40 MG/0.4 ML SYRING) | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA(CF) PEDIATRIC CROHN'S | 3 | CC | PA | QPD 0.15 per day |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | | |
|---|------|-----------------------|----|------------------|
| HUMIRA(CF) PEN (PEN 40 MG/0.4 ML, PEN 80 MG/0.8 ML) | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA(CF) PEN CROHN'S-UC-HS | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA(CF) PEN PEDIATRIC UC | 3 | CC | PA | QPD 0.15 per day |
| HUMIRA(CF) PEN PSOR-UV-ADOL HS | 3 | CC | PA | QPD 0.15 per day |
| SIMLANDI(CF) | 3 | CC | PA | QPD 0.15 per day |
| SIMLANDI(CF) AUTOINJECTOR | 3 | CC | PA | QPD 0.15 per day |

VASODILATING AGENTS

DIRECT VASODILATORS

hydralazine hcl (10 mg tablet, 25 mg tablet, 50 mg tablet, 100 mg tablet)

1

minoxidil (2.5 mg tablet, 10 mg tablet)

1

NITRATES AND NITRITES

isosorbide dinitrate

1

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS | |
|---|------|-----------------------|--------------|
| <i>isosorbide mononitrate (10 mg tablet, 20 mg tablet, 30 mg tab er 24h, 60 mg tab er 24h, 120 mg tab er 24h)</i> | 1 | | |
| NITRO-BID | 2 | | |
| <i>nitroglycerin (0.3 mg tab subl, 0.4 mg tab subl, 0.6 mg tab subl)</i> | 1 | | |
| VASODILATING AGENTS (RESPIRATORY TRACT) | | | |
| PHOSPHODIESTERASE-5 INHIBITORS (RESPIR) | | | |
| ALYQ | 3 | CC PA QPD | 2.0 per day |
| <i>sildenafil citrate 20 mg tablet</i> | 3 | CC PA QPD | 12.0 per day |
| <i>tadalafil 20 mg tablet</i> | 3 | CC PA QPD | 2.0 per day |
| PROSTACYCLIN & PROSTACYCLIN DERIVATIVES | | | |
| VENTAVIS | 3 | CC PA QPD | 9.0 per day |
| VITAMINS | | | |
| MULTIVITAMIN PREPARATIONS | | | |
| <i>prenatal with folic acid</i> | 2 | \$0 | |
| VITAMIN B COMPLEX | | | |
| <i>cyanocobalamin (vitamin b-12)</i> | 1 | QL | 1 / 30 days |

| PRODUCT DESCRIPTION | TIER | LIMITS & RESTRICTIONS |
|---|------|-----------------------|
| DIALYVITE | 2 | |
| <i>folic acid 1 mg tablet</i> | 1 | \$0 |
| MYNEPHROCAPS | 1 | |
| MYNEPHRON | 1 | |
| RENA-VITE RX | 1 | |
| RENO CAPS | 1 | |
| TRIPHROCAPS | 1 | |
| VIRT-CAPS | 1 | |
| WESCAPS | 1 | |
| VITAMIN D | | |
| <i>calcitriol (0.25 mcg capsule, 0.5 mcg capsule)</i> | 1 | |
| <i>ergocalciferol (vitamin d2) 1250 mcg capsule</i> | 1 | \$0 |
| VITAMIN K ACTIVITY | | |
| <i>phytonadione (vit k1) 5 mg tablet</i> | 1 | |

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