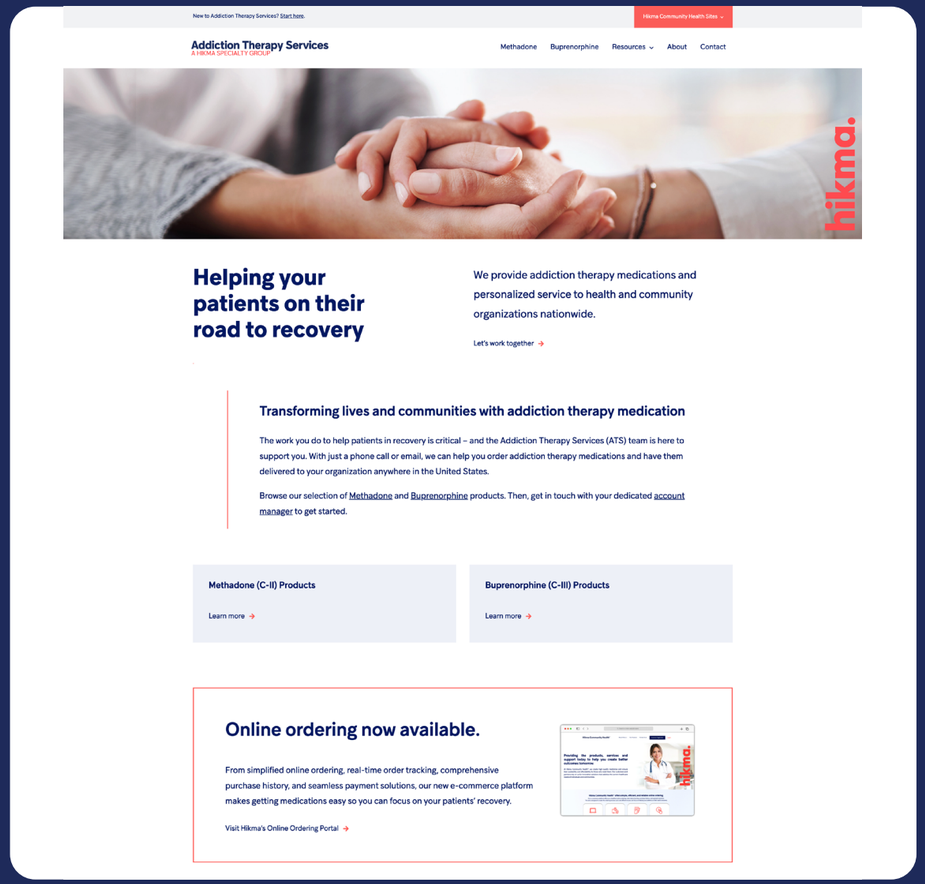
Addiction Therapy Services Product Catalog

AddictionTherapyServices.com



Addiction Therapy Services Product Catalog

Contents

Corporate Headquarters
Hikma Pharmaceuticals USA Inc.
200 Connell Drive, Suite 4000
Berkeley Heights, NJ 07922
Tel: 732.542.1191 | Fax: 732.542.0940
hikma.com/us

Addiction Therapy Services Customer Service
Business Hours: 8am ET – 5pm ET, Monday – Friday
Tel: 833.449.3475 | Fax: 866.464.1562
Email: usaddictiontherapy@hikma.com

Addiction Therapy Services Products

METHADONE HCl Oral Concentrate USP C-II

Product Description	NDC Number	Strength	Unit Size	Prescribing Information
Clear/Colorless Solution Flavorless	0054-0391-68	10 mg / mL	1000 mL Bottle	
Clear/Red Colored Solution Cherry flavored	0054-0392-68	10 mg / mL	1000 mL Bottle	

METHADONE HCl Tablets USP C-II

Product Description
Round/White Tablets
Flavorless

NDC Number	Strength	Unit Size	Prescribing Information
0054-0709-25	5 mg	Bottle - 100 tablets	
0054-0710-25	10 mg	Bottle - 100 tablets	

DISKETS® Dispersible Tablets C-II (Methadone Hydrochloride Tablets for Oral Suspension USP)

Product Description
Light Pinkish-Orange Cross-Scored Tablets
Orange-Pineapple Flavored

NDC Number	Strength	Unit Size	Prescribing Information
0054-4538-25	40 mg	Bottle - 100 tablets	

DISKETS® is a registered trademark of Hikma Pharmaceuticals USA Inc.

Addiction Therapy Services Products

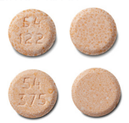
BUPRENORPHINE Sublingual Tablets C-III

Product Description
Round/White Sublingual Tablets
Flavorless

NDC Number	Strength	Unit Size	Prescribing Information
0054-0176-13	2 mg	Bottle - 30 tablets	
0054-0177-13	8 mg	Bottle - 30 tablets	

BUPRENORPHINE AND NALOXONE Sublingual Tablets USP C-III

Product Description
Round/Speckled Peach to Peach Sublingual Tablets
Lemon-Lime Flavor



NDC Number	Strength	Unit Size	Prescribing Information
0054-0188-13	2 mg / 0 .5 mg	Bottle - 30 tablets	
0054-0189-13	8 mg / 2 mg	Bottle - 30 tablets	

BUPRENORPHINE AND NALOXONE Sublingual Film C-III

Product Description
Orange/Rectangular Film
Lemon-Lime Flavor

NDC Number	Strength	Unit Size	Prescribing Information
43598-579-30	2 mg / 0 .5 mg	30 Pouches containing 1 Sublingual Film	
43598-580-30	4 mg / 1 mg	30 Pouches containing 1 Sublingual Film	
43598-582-30	8 mg / 2 mg	30 Pouches containing 1 Sublingual Film	
43598-581-30	12 mg / 3 mg	30 Pouches containing 1 Sublingual Film	

BUPRENORPHINE AND NALOXONE Sublingual Film C-III is manufactured by Lohmann Therapy Systems, (LTS), West Caldwell, NJ 07006 for Dr. Reddy's Laboratories Inc., Princeton, NJ 08540, USA. Please visit https://dailymed.nlm.nih.gov/ for additional information specific to this product, including the Full Prescribing Information with complete indications for use, warnings, precautions and adverse reactions.

Product images may not reflect actual sizes and/or exact colors.

Please visit addictiontherapyservices.com for additional product information, including the Full Prescribing Information with complete indications for use, warnings, precautions and adverse reactions for each product.

usaddictiontherapy@hikma.com | Customer Service 833.449.3475

Ordering Instructions

Mandatory items for order processing

The following must be on file with Hikma:

- DEA Registration Certificate
- State License
- Completed Due Diligence Questionnaire

C-II Ordering

Electronic Ordering

- If you are enrolled in the DEA's Controlled Substance Ordering System (CSOS), visit order.hikmacommunityhealth.com to place your order.

Paper DEA Order Form 222

- Follow directions printed on the back of the DEA-222 Form.
- Shipment can only be made to the address shown on the form.
- If your name and address have changed, contact your regional DEA office for a new supply of forms.
- 222 Form MUST be mailed. 222 Forms that are emailed or faxed will NOT be accepted. 222 Forms must be mailed to the following address:

  Hikma Pharmaceuticals USA Inc.
  1809 Wilson Rd., 08-118
  Columbus, OH 43228

- Federal Regulations do not allow us to accept forms that have corrections, alterations or write overs.

C-III Ordering

- Visit order.hikmacommunityhealth.com to place your order.

Shipping

- Our distribution center ships orders Monday through Friday.
- Next business day delivery by 10:30 to most U.S. addresses via FedEx Priority Overnight.

Hikma Customer Portal

Offering simple, efficient, and reliable online ordering

Get streamlined access to addiction therapy medication with online ordering, order status tracking, purchase history and payment solutions. We make the process quick and efficient so you can focus on what matters most—helping your patients on the road to recovery.

Order

Save time with simplified online ordering through our convenient portal.

Track

Get real-time updates as we process and ship your order.

Receive

Manage the receipt of your order with our easy-to-use tools and resources.

Pay

Easily view and pay invoices through your online payment portal.

How to Get Started

If you are an existing customer, visit order.hikmacommunityhealth.com to create your username and follow the steps below:

1. Click on **Customer Registration**.
2. Select **My facility is a Hikma customer and I need to create a username**.
3. Enter your information. Process will require your Hikma Sold To account number.
4. You will receive an email confirmation when your user account has been established.

To learn more about the Hikma Customer Portal, visit order.hikmacommunityhealth.com

Controlled Substance Ordering System (CSOS)

Benefits of CSOS

- Improves accuracy by eliminating the paper 222 form
- Quicker turn-around times
- Lower costs per transaction
- Introduces greater security and helps ensure DEA compliance

Overview

- Customer applies for a digital signing certificate issued by the DEA
- Upon approval by the DEA, customer downloads the digital certificate
- Contact Hikma to gain access to the order.hikmacommunityhealth.com website
- Customer can begin ordering their products electronically

How to apply for a digital certificate

To obtain a digital certificate, a DEA registrant must enroll in the CSOS program through the DEA website (deaecom.gov). Every individual who wants to sign electronic orders for controlled substances must enroll with the DEA to acquire his or her own personal digital certificate.

Choose your applicant type

- Registrant – Individual who is authorized to sign DEA applications. The Registrant applicant must name him/herself as Coordinator or delegate the role to another Principle Coordinator or existing subscriber.
- Coordinator – Any individual employed by the DEA Registrants organization. The Principle Coordinator is a required role for each DEA registration number in the CSOS program and will be the primary CSOS contact. If the Registrant is requesting to be the Coordinator, he or she should apply only as a registrant and indicate him or herself as the Coordinator.
- Power of Attorney – Individual with the authority to sign controlled substance orders for a DEA Registrant.

What happens after I enroll

Once the enrollment process is complete, the DEA will send an activation notice via email and postal mail containing your access code and password. These two items (access code and password) must be married up in order for you to retrieve your digital certificate from the DEA website.

When all documentation has been received, contact Hikma Addiction Therapy Services (833.449.3475) to gain access to the order.hikmacommunityhealth.com website. Hikma will assist you in setting up your account and provide training for use of the system.

C-II Ordering Instructions

Sample Single-Sheet 222 Form with instructions

PURCHASER INFORMATION

REGISTRATION INFORMATION
REGISTRATION #:
REGISTERED AS:
SCHEDULES:
ORDER FORM NUMBER:
DATE ISSUED:

SUPPLIER DEA NUMBER:# RW0498419 (2)
PART 2: TO BE FILLED IN BY PURCHASER
Hikma Pharmaceuticals USA Inc.
BUSINESS NAME
1809 Wilson Road, 08-18
STREET ADDRESS
Columbus, OH 43228
CITY, STATE, ZIP CODE

PART 1: TO BE FILLED IN BY PURCHASER
Name and Title
Print or Type Name and Title
Sign and indicate Signature Authority — Current Date
Signature of Requesting Official (must be authorized to sign order form) — Date

PART 3: ALTERNATE SUPPLIER IDENTIFICATION - to be filled in by first supplier (name in part 2) if order is endorsed to another supplier to fill. (3)
ALTERNATE DEA #
Signature- by first supplier
OFFICIAL AUTHORIZED TO EXECUTE ON BEHALF OF SUPPLIER — DATE

PART 5: TO BE FILLED IN BY PURCHASER (5)

ITEM	NO. OF PACKAGES	PACKAGE SIZE	NAME OF ITEM	NUMBER REC'D	DATE REC'D	PART 4: TO BE FILLED IN BY SUPPLIER NATIONAL DRUG CODE	NUMBER SHIPPED	DATE SHIPPED
1	100	1000mL	Methadone HCl Oral Conc- Cherry					
2	80	1000mL	Methadone HCl Oral Conc- Clear					
3	50	100	Methadone HCl 40mg Diskets					
4	20	100	Methadone HCl Tablets, 5mg					
5	10	100	Methadone HCl Tablets, 10mg					(4)
6								
7								
8								
9								
10								
11								
12								
13								
14								
15								
16								
17								
18								
19								
20								
	LAST LINE COMPLETED (MUST BE 20 OR LESS)							

(1)

1 Purchaser Information:
- Print/Type name and title.
- Sign and indicate signature authority. For example, "attorney-in-fact", "power of attorney", "designated agent", or "secretary" may be used.
- Fill in current date.
- Customer must complete the following columns:
  - No. of packages
  - Package size
  - Name of item(s): Please distinguish between Cherry and Clear Methadone HCl Oral Conc.
  - Fill in "Last Line Completed"

2 Supplier Identification
- Enter DEA number, name, and address of supplier.
  DEA#: RW0498419
  Hikma Pharmaceuticals USA Inc.
  1809 Wilson Road, 08-118
  Columbus, OH 43228

3 Alternate Supplier Identification
- Leave this portion of the form blank.

4 Controlled Substance Shipment
- Leave this portion of the form blank.

5 Controlled Substance Receipt
- Purchaser will enter the number of packages received and date received for each line item on its copy of the original order form.

Purchaser must make a copy of the original order form for its records before mailing the original to the supplier.

Mail the original copy to the address below.
Hikma Pharmaceuticals USA Inc.
1809 Wilson Road, 08-118
Columbus, OH 43228

Return Goods Policy Effective January 1, 2025

Hikma Pharmaceuticals USA Inc. ("Hikma") Return Goods Policy (this "Policy") applies to all Hikma labeled pharmaceutical products ("Products") manufactured and/or distributed in the United States and its territories, commonwealths, and possessions ("Territory"). This Policy applies to Products from direct customers and distributors of Hikma, and indirect customers returning through the wholesaler pursuant to the original purchase from Hikma ("Customers"). Unless otherwise required by regulations, laws, or expressly agreed upon by the parties, this Policy applies to Products.

CLAIM PREREQUISITES

- In the specific event of either an overage, shortage, concealed shortage, damage, or incomplete shipment of Product ("Claims"), Customer shall contact Hikma's Claims department within one (1) business day of identification of issue(s) and coordinate with Hikma on commercially reasonable efforts to reconcile shipment errors, including Transaction Information/Transaction Statement (TI/TS) data pursuant to the Drug Supply Chain Security Act (DSCSA).
- Claims may be denied in situations wherein Customer has not conducted commercially reasonable efforts to provide supporting documentation (e.g., serial numbers) to Hikma for the required Transaction Information/Transaction Statement (TI/TS) data.
- In the event of an EPCIS file failure, Customer shall contact Hikma's serialization department within one (1) business day of identification of issue(s) and coordinate with Hikma on commercially reasonable efforts to reconcile errors, including Transaction Information/ Transaction Statement (TI/TS) data prior to initiating any Request for a Return Authorization ("RA").

RETURN AUTHORIZATION PROCEDURE FOR EXPIRED PRODUCTS

- RA and box labels may be made by any of the below methods through Hikma's third-party reverse logistics processor, Inmar Intelligence ("Inmar"):
  1. Visit Inmar's website at: https://returns.healthcare.inmar.com.
     - An uploaded PDF copy of your debit memo is required.
  2. E-mail the debit memo to: rarequest@inmar.com.
     - Must include: NDC#, Lot#, Expiration Date, and Unit Price for each item being returned.
  3. Fax your debit memo to Inmar at: 817-868-5343.
     - All third-party return processors must contact Inmar for a RA using one of the above methods.
     - Upon receipt of a RA and box labels, actual returns are to be forwarded to the processing facility at the following location:
       Inmar Intelligence
       3845 Grand Lakes Way, Suite 125, Grand Prairie, Texas 75050
- Questions related to returns of expired Products may be sent to: expiredreturns@hikma.com

RETURN AUTHORIZATION PROCEDURE FOR NON-EXPIRED RETURNS

- "Non-Expired Returns" is defined as the return of Product for any reason other than expiration including Claims as defined herein.
- Any Claims must be adjudicated and resolved prior to receiving a RA.
- Email Hikma's Claims Department at: usclaims@hikma.com.
  - Include: NDC#(s), Lot #(s), Serial Number(s), Purchase Order Number, Quantity.
- For damaged Product, photos must be submitted.
- Hikma will send a RA, box label(s), and Call Tag(s).
  - Upon receipt, Products are to be forwarded to the processing facility as indicated on the RA.
- If Products are C-II controlled substances ("Controlled Products"), you will receive a DEA Form 222 from Inmar or Hikma.
  - DEA Form 222 must be included with Controlled Products.

RETURN AUTHORIZATION PROCEDURE FOR RECALLED PRODUCTS

- For Recalled Product or market withdrawal Product, please refer to your directions as indicated on your Recall Response Form.
- Questions related to Recalled Products for Hikma can be sent to: usrecall@hikma.com.

PRODUCT RETURNS ELIGIBLE FOR REIMBURSEMENT

Products eligible for reimbursement include the following:

- Authorized Expired Product, which is Product returned in full and unopened containers with a Hikma label, purchased directly from Hikma and returned directly to Inmar: (i) within six (6) months prior to; or (ii) within twelve (12) months after the expiration date.
- Recalled Product, as stated on a recall notice issued by Hikma, which is returned directly to Inmar after requesting and receiving an RA from Inmar.
- Products which are authorized Non-Expired Returns, purchased from Hikma, and returned due to Claims.

PRODUCT RETURNS INELIGIBLE FOR REIMBURSEMENT

- Products ineligible for reimbursement include the following:
- Product(s) returned: (i) earlier than six (6) months prior to the expiration date; or (ii) greater than twelve (12) months after the expiration date assigned to such Product.
- Partial units or containers, except where mandated by federal, state, or local laws.
- Product(s) not in their original, sealed, full, unopened, and unadulterated Hikma container including an inner pack, unit or vial with a non-saleable NDC.
- Private labelled, re-packaged, re-constituted, and/or contract manufactured Product(s).
- Product(s) sold by Hikma at no cost including, but not limited to, donations and samples.
- Product(s) sold as short-dated, close-out, special promotion, and/or sold as non-returnable.
- Product(s) not purchased directly from Hikma or the Customer's authorized distributor/wholesaler.
- Product(s) returned by an indirect Customer for which the distributor/wholesaler did not purchase the listed Product NDC and Lot# and/or Serial Number from Hikma.
- Product(s) with defaced or missing Hikma labels which do not clearly display the Product's expiration date, NDC, and/or valid Lot number.
- Product(s) damaged or deteriorated due to: (i) negligence; (ii) improper handling or storage by the Customer; or (iii) insurable causes such as fire, floods, and/or natural disasters.
- Customer overstocked Product(s) unless prior written approval from Hikma is received.
- Product(s) sold by Hikma Injectables Inc. d/b/a Hikma 503B which are not eligible for credit per Hikma 503B's Return Goods Policy.
- Product(s) purchased through a bankruptcy sale.
- Product(s) received by Inmar thirty (30) days or more after the date assigned on the RA.
- Product(s) purchased or distributed contrary to federal, state, or local laws.
- Product(s) sold directly by Hikma or through an authorized wholesaler or distributor of record to any city/municipality, county, state, and/or federal entity for the purpose of stockpiling.
- Product(s) purchased outside of the Territory.
- Product(s) purchased for future events including speculative purposes.
- Products destroyed and/or not received by Inmar, unless approved in writing by Hikma.

- Expired Product(s) with a returnable credit value of $25.00 or less per debit memo, as determined by Hikma's pricing valuation.
- Expired Product(s) whose cumulative calendar year credit value has exceeded a one (1%) percent limitation based on Customer's prior calendar year's purchase value of all return eligible Products. Return value based on return credit dollars issued and includes both direct and indirect/third party Customer expired returns.

VALUATION OF EXPIRED RETURNS CREDIT MEMOS

- For direct Customers, a credit will be issued based upon the lower of the current net invoice price at the time the Product(s) is received by Inmar -OR- the lowest net invoice price in the prevailing twenty-four (24) months -OR- lowest actual net price if able to be determined by Lot number and/or Serial number.
- For indirect Customers, a credit will be issued based upon the lower of the current net indirect price at the time the Products are received by Inmar -OR- the lowest net indirect contract price ("LNICP") in the prevailing twenty-four (24) months from the wholesaler. If Hikma cannot identify the LNICP for a Customer, then Hikma will use a predetermined indirect return price.
- Indirect returns will be credited to the wholesaler or distributor of purchase.

VALUATION OF NON-EXPIRED RETURNS CREDIT MEMOS

- For Non-Expired Returns, including but not limited to Claims, a credit will be issued based on the net invoice price of the Product(s) as purchased.
- Hikma may reduce the credit value with a restocking fee of 20% of the net invoice price of the Product(s) if the cause of the return is due to no fault of Hikma.
- For recalled Product, current net sale price will be credited.

CREDIT MEMO CONDITIONS

- The amount of credit issued or authorized by Hikma is directly correlated to the quantity validated by Inmar or Hikma. In the event of any conflict between the Customer's claimed quantity and the quantity validated by Inmar or Hikma, the quantity validated by Inmar shall control.
- Credit will be issued by Hikma in the form of a credit memo only.
- Customer deductions for returns must reference Hikma's issued Credit Memo number or the Debit Memo number as supplied to Inmar.

SHIPPING ERRORS

- Hikma must be notified of any shipping disputes within three (3) business days of receipt of Product(s). Product(s) shipped in error by Hikma must be returned within thirty (30) days of shipment to receive credit. Product(s) returned after thirty (30) days will not be eligible for credit.
- If a shipping error involves any Controlled Products, Hikma must be notified within 24 hours of receipt of the order of any overages, shortages, or mistakes in such Controlled Products order.
- For non-Controlled Products, a Customer will make best efforts to retain an overage shipment and the Customer and Hikma shall mutually agree on the pricing for such manually processed invoice and related needed data including but not limited to ASN.

THIRD PARTY PROCESSORS

- Third party processors and reverse logistics companies must comply with all requirements of this Policy. Hikma will not pay or reimburse any service fees to the purchaser or third-party return processor, including handling fees, processing fees, or freight charges incurred.
- Hikma will not process returns using pricing from any third party's internally generated price list. Pricing will be based on Hikma' valuation as described in this Policy.

TRANSPORTATION

- Transportation charges, including prepaid freight and insurance, are the responsibility of the customer except when due to a Hikma error, as determined by Hikma.
- Hikma is not responsible for lost or damaged shipments of Product(s). Insuring and tracking shipments are the responsibility of the customer.

COMPANY DISCLAIMERS

- Submission of Product does not constitute Hikma's acceptance for credit.
- Package size, Lot number and Lot expiration date will be obtained and verified after receipt of Product by Inmar.
- Returns are subject to review by Hikma and issuance of an RA number does not guarantee credit.
- Hikma reserves the sole right to determine whether Products qualify for return or credit.
- Inmar's determination of the physical count of Products will be final. By returning Products you authorize Hikma and its designee, as your agent, to destroy, without payment or other recourse.
- Any and all credits provided pursuant to this Policy are only valid if redeemed within one (1) year of issuance. Any and all credits that are not redeemed within one (1) year of issuance shall be null and void, except where not permitted by state or federal laws.
- Customers will ensure that a debit memo claim and Products are received within fifteen (15) calendar days of the debit memo date by Hikma or Inmar to receive credit.
- Unauthorized deductions for Product(s) will not be accepted.
- Hikma reserves the right to require proof of purchase source on all Products returned for credit/refund.
- Non-Hikma product(s) returned will not be the responsibility of Hikma. Hikma reserves the right to charge customers for any costs incurred to process and destroy such non-Hikma product(s). Any such non-Hikma product(s) will not be returned to the customer.
- Serial numbers verified by Inmar and/or Hikma that do not correspond to the customer submitting the return or the customer returning the Product may be denied for credit.

This Policy supersedes all previous policies and may be modified or updated by Hikma at its discretion. Hikma values the relationship it shares with its customers and will make a commercially reasonable attempt to provide thirty (30) days advance notification of any change to this Policy.

Customers will be expected to adhere to the most current policy which can be found on the Hikma website: www.hikma.com.

Addiction Therapy Services
833.449.3475
usaddictiontherapy@hikma.com
addictiontherapyservices.com

200 Connell Drive, Suite 4000, Berkeley Heights, NJ 07922
Customer Service 833.449.3475 | hikma.com/us
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