Indiana Department of Health-Immunization Division



2023 VACCINE MANAGEMENT PLAN

OFFICE AND VACCINE COORDINATOR INFORMATION

This plan must be reviewed and updated annually (every 12 months) or when key staff change (primary/back-up coordinators and/or signatory). This plan must be dated and signed by the person responsible for the VMP content.

Office/Practice Name			
VFC PIN (6 Digits- e.g. M00L00)			
Physical Address			
Phone Number			
Fax Number			
			•
	Name:		
Medical Director/Signatory	Email:		
	Contact Number:		
	T		
	Name:		
Primary Vaccine Coordinator	Email:		
	Contact Number:		
	T		
	Name:		
Back-Up Coordinator	Email:		
	Contact Number:		
			
Person(s) with 24-hour access to	Name:		
building, in case of emergency	Contact Number:		
Reviewed on	by		
Date		Signature of person filling	
Juli	-	out this form	

Indiana Department of Health-Immunization Division



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SIGNED PROVIDER AGREEMENT PAGE (COPY)

Place a copy of your office's signed provider agreement to receive publicly supplied vaccine here

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VACCINE COORDINATOR ROLES AND RESPONSIBILITIES

Summary

VFC providers are required to designate a primary vaccine coordinator and at least one back-up vaccine coordinator for each facility. Notify the IDOH Regional Quality Assurance Specialist immediately of any changes in the medical director, primary vaccine coordinator, or back-up vaccine coordinators via the email or phone number listed below. All coordinators must be fully trained on routine and emergency standard operating procedures (SOPs) for vaccine ordering, storage, handling, transport, and inventory management.

The vaccine coordinator is responsible for overseeing all vaccine management within the facility, including:

- Developing and maintaining this Vaccine Management Plan
- Monitoring all vaccine storage and handling and vaccine administration practices in the facility, including
 responding to and documenting all temperature excursions and reporting them to the Indiana Department
 of Health (IDOH) Staff Representative
- Overseeing vaccine ordering and notifying the IDOH Staff Representative if vaccines will expire before they
 are administered
- Ensuring and documenting annual vaccine management training for designated staff as well as training new staff upon hire
- Participating in and documenting completion of annual training on VFC requirements
- Storing all required documentation for three years

IDOH Regional Quality Assurance Specialist	
Contact Number	
Email	
Immunization Division Central Office Contact	800-701-0704 vaccine@health.in.gov (always include your VFC PIN for reference)

Please reference the map on the next page, to identify the Regional Quality Assurance Specialist for your facility.

Key Personnel

Maternal Immunization and Perinatal Hepatitis B Coordinator:

Ondreya Witmer 317-518-0726 owitmer@health.in.gov

ROAS Coordinator:

Diana Taylor, 317-416-5929 ditaylor@health.in.gov

VAM Coordinator:

Tracy Brunette, 317-606-5091 tbrunette@health.in.gov

North A

RQAS:

Cynthia Joiner, 317-967-5956 cjoiner@health.in.gov

VAM:

Melissa Sailors,317-376-9783 msailors@health.in.gov

North B

ROAS:

Mindy Dixson, 317-914-5286 mdixson2@health.in.gov

VAM:

Melissa Sailors,317-376-9783 msailors@health.in.gov

Central C

RQAS:

Ashlei VanBuskirk 463-271-1238 avanbuskirk@health.in.gov

VAM:

Nancy Fetsch, 317-233-2320 nfetsch@health.in.gov

Central D

RQAS:

Carla Cly-Williams, 317-439-6370 cclywilliams@health.in.gov

VAM:

Nancy Fetsch, 317-233-2320 nfetsch@health.in.gov

South E

RQAS:

Tom Keller, 317-519-3052 tkeller@health.in.gov

VAM:

Holly Carson, 317-726-7426 hcarson@health.in.gov

South F

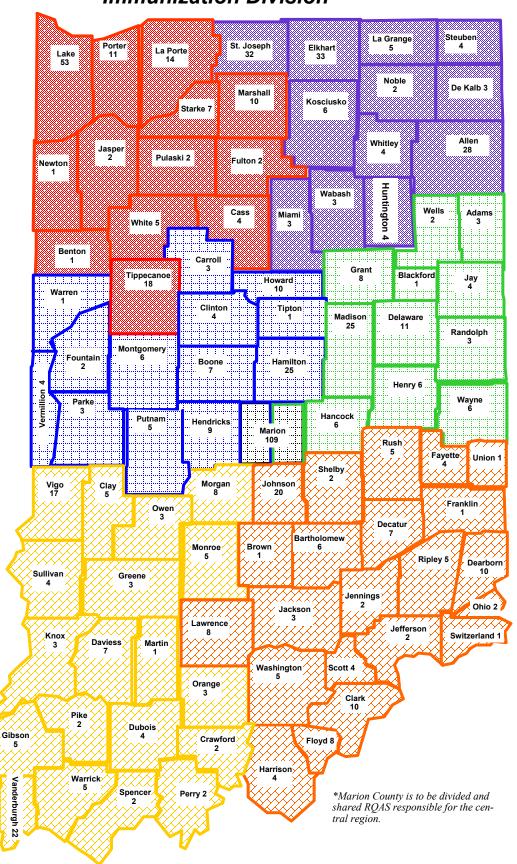
ROAS:

Milly Jines, 317-519-2611 mjines@health.in.gov

VAM:

Holly Carson, 317-726-7426 hcarson@health.in.gov

Indiana State Department of Health Immunization Division



Regional Quality Assurance Specialists (**RQAS**)
Vaccine Ordering and Accountability (**VAM**) Specialists
Immunization Division Service Delivery Map, December 2022

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More specifically, vaccine coordinators are responsible for the following:

Vaccine Storage and Handling

- Ensure the vaccine cold chain is maintained at all times.
- Ensure refrigerator and freezer units that house VFC vaccine meet CDC requirements.
- Ensure storage unit digital data loggers (DDLs) and/or continuous temperature monitoring system meet CDC requirements.
- Oversee proper receipt and storage of all vaccine shipments, including training any office and front desk staff on storage and handling practices in the event they encounter vaccine shipments.
- Monitor all VFC inventory expiration dates and rotate vaccine stocks to ensure soon-to-expire vaccines are used first.
- Ensure VFC and privately purchased vaccine stocks are clearly separated.
 - Have a tracking system in place to record any doses that are accidentally borrowed from VFC and ensure all VFC doses are replaced with private-purchased vaccines in a timely manner.
- Place "DO NOT UNPLUG" stickers by the wall outlets into which storage units that house VFC vaccine are plugged.
- Circuit Breaker boxes should be clearly marked with the associated outlet and emergency contact information if the breaker box is tripped and/or there is scheduled maintenance.

Vaccine Ordering and Accountability

- Submit vaccine orders via the Vaccine Ordering Management System (VOMS) in the electronic Immunization Information System (IIS) registry CHIRP (Children and Hoosiers Immunization Registry Program).
- Electronically reconcile VFC vaccine inventory monthly and submit via CHIRP.
- Account for all wasted/expired/spoiled/borrowed vaccines in CHIRP and complete/submit all associated documentation forms to the IDOH Immunization program.
- Ensure all vaccine orders are received electronically in CHIRP.

Emergency Situations

- Follow all protocols for protecting vaccines in emergency situations such as power outages, natural disasters, and equipment failure.
- Respond and manage all temperature excursion events.
- Follow all vaccine transport guidance.

VFC Programmatic Tasks

- Participate in all program compliance site visits with an IDOH Staff Representative, which may be scheduled or unannounced.
- Complete and submit annual provider recertification.

Daily Temperature Monitoring and Recording

- Document the minimum and maximum temperatures of each storage unit at the start of each workday.
- Record the current storage unit temperature twice daily on IDOH temperature logs.
- Maintain temperature monitoring devices, including a back-up device, and ensure they all have current certificates of calibration.
- Download and review DDL reports each work week, and report temps to IDOH as requested.
- Maintain storage equipment, including ensuring units are properly cleaned and serviced at all times.

Annual training requirements

 Annually, primary and back-up VFC coordinators, and any staff members that have any responsibility for VFC vaccines must complete the VFC Provider Annual Training on LMS Invest. This can be found on the CHIRP Homepage.



2023 Vaccines for Children (VFC) Vaccine Management Plan

Standard Operating Procedures (SOPs) Template

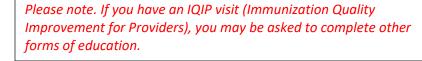
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STAFF TRAINING AND DOCUMENTATION

The IDOH Immunization Program requires primary and back-up vaccine coordinators to complete the following annually:

- VFC Provider Annual Training module in LMS INvest.
 - ► This can be found on the CHIRP Homepage.





This must be completed by any staff that will receive vaccine shipments.

It is best practice that ALL office staff complete this training.

Staff Name	LMS Training Date	Additional Education (Optional)	Additional Education (Optional)

2023 Vaccines for Children (VFC) Vaccine Management Plan

Standard Operating Procedures (SOPs) Template

Indiana Department of Health-Immunization Division



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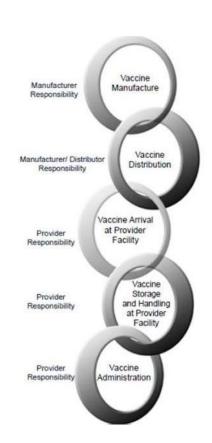


VACCINE STORAGE AND HANDLING PRACTICES

All vaccine storage and handling requirements and recommendations are in place to ensure the vaccine cold chain is maintained. The cold chain begins at the manufacturing plant, includes delivery to and storage at the provider facility, and ends with the administration of vaccine to the patient. Too much exposure to heat, cold, or light at any step in the cold chain can result in a loss of vaccine potency. Once potency is lost, it cannot be restored.

VFC Storage and Handling Requirements

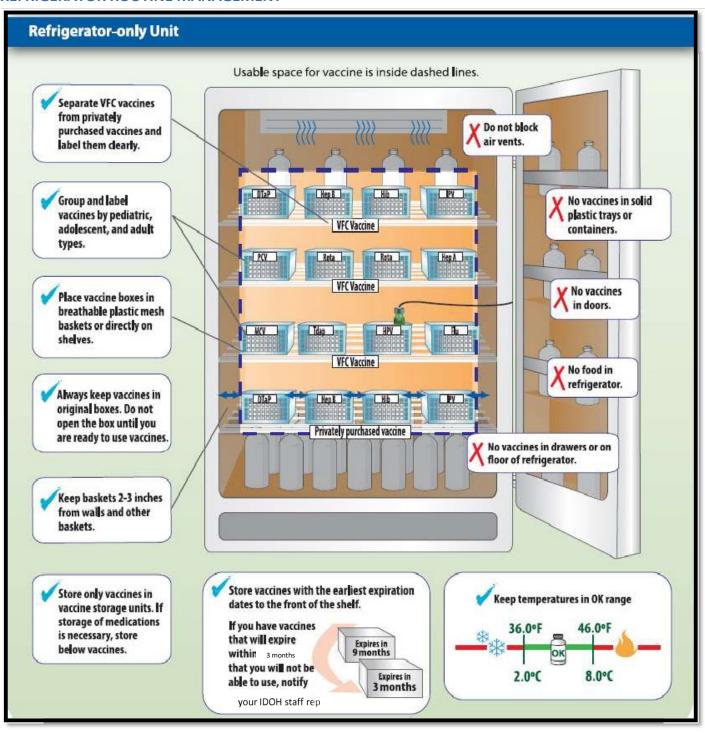
- Stand-alone storage units that maintain correct temperatures at all times.
 - Refrigerator temperature between 36°F and 46°F (2°C and 8°C); aim for 40°F (4°C)
 - Freezer temperature between -58°F and +5°F (-50°C and -15°C); aim for 0°F (-17°C)
 - Water bottles in storage units-against walls, in the back, on the floor- to help stabilize temperatures
 - Needs to have shelves for vaccine placement
 - NO food or drinks in the storage units
 - NO vaccine storage in the unit door, on the floor, and/or under the cooling vents of a storage unit
 - NO expired vaccine in unit
- Digital data loggers (DDLs) with continuous monitoring capabilities and a current, valid Certificate of Calibration Testing for each device, as well as at least one back-up DDL for each clinic.
 - Must have a glycol or other buffered solution-encased probe
 - o Probe must be placed in the central area of the storage unit
 - Record DDL minimum and maximum temperatures once daily at the start of the workday
 - Download, review, and store digital reports once a week; reset DDL memory at this time
 - Record point-in-time temperature readings twice daily on standard temperature logs
- DO NOT UNPLUG signs by the electrical outlets into which the units are plugged
- WARNING-EXPENSIVE VACCINE IN STORAGE signs on the circuit breaker with emergency contact info and the breaker numbers listed; outlet breaker connected to the units clearly marked inside the box



Indiana Department of Health-Immunization Division



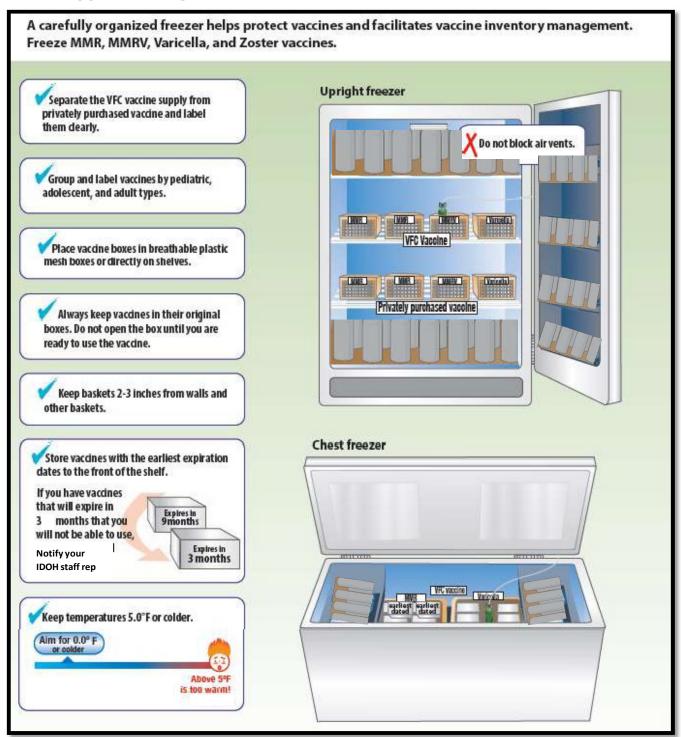
REFRIGERATOR ROUTINE MANAGEMENT



Indiana Department of Health-Immunization Division



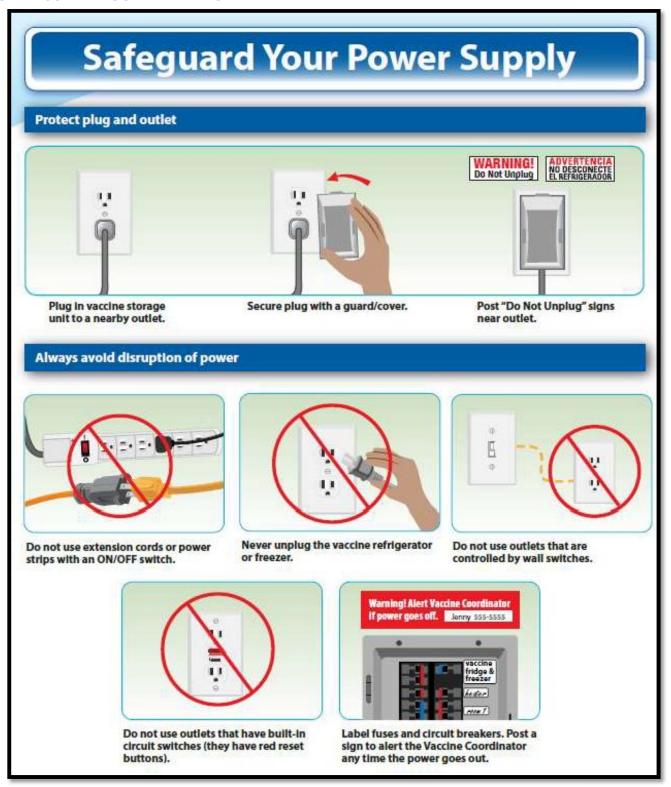
FREEZER ROUTINE MANAGEMENT



Indiana Department of Health-Immunization Division



POWER SUPPLY ROUTINE MANAGEMENT



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VACCINE STORAGE UNIT/LOCATIONS AND MAINTENANCE

Maintenance/Repair Company	Phone	

Unit Type	Unit/Location ID	Brand	Model	Dates of Repair
Refrigerator				
Freezer				

Location of completed
temperature logs
temperature logs

TEMPERATURE MONITORING DEVICE LIST AND MAINTENANCE

Calibration Company/Laboratory	Contact	Phone	
Location of Certificates of Calibration			
Location of Backup Data Logger			

Digital Data Logger or Continuous Monitoring System Device Model/Serial Number	Primary?	Backup?	Calibration Expiration Date	Alarm Setting Low	Alarm Setting High

Indiana Department of Health-Immunization Division



Place a copy of your office's current calibration certificates here.

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VACCINE ORDERING PROCEDURES

- All vaccine orders are submitted through the Vaccine Ordering Management System (VOMS) in CHIRP
 - Providers will be ordering influenza through VOMS. We have discontinued the use of paper influenza order forms.

At minimum, providers should order enough vaccine for a four-week period, but no more than three months.

- At the time of vaccine order placement:
 - Providers must ensure contact information and delivery hours are correct for the next three weeks to ensure accurate and prompt vaccine delivery (e.g. if your office will be closed on any day in the coming three weeks, or open for less than four (4) hours, uncheck that day as an option when placing your order).
 - o Providers must submit their inventory monthly in VOMS.
 - All vaccines administered must be entered into CHIRP prior to inventory reconciliation, via manual or Electronic Medical Record (EMR) entry.
- Non-frozen vaccine orders typically arrive 7 to 10 days after the order has been approved in VOMS.
- **Frozen** vaccine ships directly from the manufacturer (Merck) and will arrive separately from refrigerated vaccine. Orders may arrive up to 14 days after the VOMS approval date.

VACCINE RECEIVING AND DOCUMENTATION PROCEDURES

The primary and/or back-up coordinator must be immediately notified if any vaccine shipment arrives. Coordinators must immediately unpack, store, and document vaccines and diluents upon receipt.

Actions must include:

- Examining the shipping container and vaccine vials for signs of physical damage
- Comparing the contents of the container to the packing list to be sure they match
 - If there is a discrepancy with the order, provider must immediately notify the Indiana Department of Health-Immunization Division at 1-800-701-0704.
- Making sure lyophilized (freeze-dried) vaccines came with the correct type and quantity of diluents
- Checking both vaccine and diluent expiration dates to ensure none are expired or soon-to-expire products
- Checking the cold chain monitor (CCM) for any indication of a temperature excursion during transit
- Receiving vaccine order via VOMS so that it populates your online inventory before staff begin using it

WITHIN TWO HOURS OF VACCINE DELIVERY:

If any damage, excessive shipping time, cold chain breach is suspected notify the Indiana Immunization Program at 1-800-701-0704.

Indiana Department of Health-Immunization Division



VACCINE INVENTORY CONTROL PROCEDURES

Ensure proper vaccine handling and preparation practices

- Vaccines should be prepared immediately prior to administration.
- Reconstitute lyophilized vaccine with the diluent that came with the vaccine- nothing else.
- Always check expiration dates prior to preparing the vaccine (and its diluent).
- A single-dose vial contains one dose and should only be used for one patient.
- A multi-dose vial is intended for ten (10) doses only and should not be punctured more than ten (10) times.

o In the event of a "drawn but not used" vaccine:

- Label the vaccine with type, lot number, date/time prepared, and initials of the preparer, and place into the appropriate storage unit (refrigerator or freezer).
- Discard any refrigerated, pre-drawn doses no later than the end of the workday.
- Varicella-containing products must be used within 30 minutes of reconstitution.

Monitor multi-dose vial use

IPV multi-dose vials can be used to their expiration date and do not have to be discarded after 28 days of opening.

Monitor vaccine inventory expiration dates

- With each vaccine shipment, organize storage unit so that vaccine with the closest expiration date is placed in front to be used first.
- o Conduct a patient reminder/recall for vaccine types that are soon-to-expire.
- If you have vaccine that will expire within 90 days (three months) that your office does not
 anticipate using, and you have conducted a reminder/recall, notify your IDOH staff representative
 to inquire about a possible transfer to another VFC provider.

Minimize VFC vaccine borrowing

- VFC vaccine <u>cannot</u> be used as a replacement system for a provider's privately purchased vaccine inventory.
- Borrowing between inventory stocks should be rare.
- Documentation must occur when any vaccine is borrowed regardless of inventory origin (e.g. VFC to Private or Private to VFC).
- The Borrowing Report is the only form of acceptable documentation and it must specify an
 approved reason for vaccine borrowing. The form must be submitted within 30 days of borrowing
 via email vaccine@health.in.gov or fax 317-233-3719.

Approved borrowing reasons:

- Vaccine shipment delay (public or private)
- Vaccine not useable on arrival (public or private)
- Ran out of vaccine between orders [not due to shipping delays] (public or private)
- Short-dated vaccine from one stock used on a patient with the opposite eligibility (public or private)
- Accidental use of vaccine from one stock on a patient with the opposite eligibility (public or private)
- Replacement of private dose with VFC when insurance plan did not cover vaccine (public)

Indiana Department of Health-Immunization Division



- When borrowing occurs, funding source integrity needs to be maintained both when the vaccine is originally borrowed and when stock is repaid.
- The funding sources for borrowed and replaced doses will need to be adjusted in VOMS.
 - Quick reference guides are located on the IDOH website https://www.in.gov/health/immunization/.
- Please note: providers may use private stock seasonal influenza vaccine to vaccinate VFC-eligible children if VFC seasonal influenza stock is not yet available. Those private stock doses used on VFCeligible children can later be replaced when VFC stock becomes available. This one-directional borrowing exception is unique to seasonal influenza vaccine.

VACCINE LOSS (EXPIRED/WASTED/SPOILED) MANAGEMENT PROCEDURES

All VFC providers are required to document and report all incidents of vaccine loss and wastage. When managing expired, spoiled, and wasted vaccine, providers must:

- Remove the vaccines from any storage unit that stores viable vaccines.
- Label vaccines DO NOT USE.
- Document wasted or expired vaccines in VOMS.
 - All returns must be completed and documented in VOMS within six (6) months after the product expiration or waste date.

Expired vaccine

- ALL expired vaccine must be returned.
- Primary vaccine coordinators will receive a McKesson return label via email within thirty (30) days of reporting the return in VOMS.

WHAT SHOULD <u>NOT</u> BE RETURNED TO MCKESSON

- Used syringes, with or without needles
- Broken vials
- Wasted products such as a syringe that was drawn up but not used
- Any multi-dose vial from which some doses have been withdrawn
- Diluent (expired or not expired)
- IG, HBIG, PPD
- Private-purchased vaccine

Discard these items according to your facility's procedures.



VFC ELIGIBILITY AND INSURANCE

*Patients must be screened for VFC eligibility at each visit

Eligibility for Publicly Funded Vaccines Reference Guide for:

Vaccines for Children (VFC) providers, Adults Vaccine Program (AVP) providers, and Electronic Medical Record contacts.

Patient Status (<19 years old)	Eligible for Public Vaccine	IIS Eligibility Category Selection	IIS Eligibility Code & Description	IIS Funding Source and Description	Billing
American Indian/ Alaskan Native • Child is less than 19 years • As defined by the Indian Health Care Improvement Act (25 U.S.C 1603-13)	Yes Federal (VFC) vaccine eligible	American Indian/ Alaskan Native	V04 VFC eligible - American Indian/ Alaskan Native	VXCI - Publicly funded vaccine stock	 Cannot bill for the cost of the vaccine Can bill administration fee up to \$20.32 per dose May issue only a single bill within 90 days Cannot send to collections for unpaid administration fees
 Uninsured Child is less than 19 years Does not have health insurance Enrolled in a Health Care Sharing Ministry (i.e., MediShare, Liberty Healthshare, HealthShare Altura, Christian Healthcare Ministries, Solidarity Healthshare) 	Yes Federal (VFC) vaccine eligible	Uninsured	V03 VFC eligible - Uninsured	VXCI - Publicly funded vaccine stock	 Cannot bill for the cost of the vaccine Can bill administration fee up to \$20.32 per dose May issue only a single bill within 90 days Cannot send to collections for unpaid administration fees
 Underinsured Child Child is less than 19 years Has insurance that does not cover vaccine Served at a Federally Qualified Health Center/ Rural Health Center (FQHC/RHC) or deputized Local Health Department (LHD) 	Yes Federal (VFC) vaccine eligible	Underinsured	V05 VFC eligible - Underinsured	VXCI - Publicly funded vaccine stock	 Cannot bill for the cost of the vaccine Can bill administration fee up to \$20.32 per dose May issue only a single bill within 90 days Cannot send to collections for unpaid administration fees
Other • Child is less than 19 years • Has insurance that does not cover vaccine • Served at any non-FQHC or LHD VFC provider location	Yes Federal (VFC) vaccine eligible	Underinsured	VOS VFC eligible - Underinsured	VXCI - Publicly funded vaccine stock	 Cannot bill for the cost of the vaccine Can bill administration fee up to \$20.32 per dose May issue only a single bill within 90 days Cannot send to collections for unpaid administration fees

2023 Vaccines for Children (VFC) Eligibility Guide

Indiana Department of Health-Immunization Division



Indiana Family Services (FSSA) offers several different programs and services under the Indiana Health Coverage Programs (IHCP). Each IHCP member is issued a number referred to as the Member ID assigned by FSSA Division of Family Resources (DFR). They type of card received depends on the IHCP program in which the member is enrolled. Providers are required to verify member eligibility on the date of service via the Provider Healthcare portal. Providers that fail to verify eligibility are at risk of claims being denied due to membership ineligibility or coverage limitations.

Viewing a member ID card alone does not ensure member eligibility.

- Hoosier Health Cards are issued by FSSA DFR.
- Hoosier Care Connect members receive member ID cards from their individuals
 Managed Care Entities (MCEs): Anthem or MHS
- Hoosier Healthwise members receive member ID cards from their individual MCEs: Anthem, CareSource, MHS, and MDwise.
- Package A (Medicaid) is not listed on the card and can only be determined via the Provider Healthcare Portal.

Patient Status (<19 years old)	Eligible for Public Vaccine	IIS Eligibility Category Selection	IIS Eligibility Code & Description	IIS Funding Source and Description	Billing
 Medicaid Child is less than 19 years old Enrolled in Traditional Medicaid Enrolled in Hoosier CareConnect (Full Medicaid of Package A) Enrolled in Hoosier Healthwise Package A Package A is not listed on the card and can only be determined via the Provider Healthcare Portal. 	Yes Federal (VFC) vaccine eligible	Medicaid	V02 VFC eligible – Medicaid	VXC1- Publicly funded vaccine stock	 Cannot bill for the cost of the vaccine Can bill administration fee up to \$20.32 per vaccine dose May issue only a single bill within 90 days Cannot send to collections for unpaid administration fee
Private Insurance • Child is less than 19 years old • Enrolled in private insurance plan	No	Ineligible for VFC/ Private Insurance	V01 Ineligible for VFC/ Private Insurance	РНС70	Bill according to plan guidelines

2023 Vaccines for Children (VFC) Eligibility Guide

Indiana Department of Health-Immunization Division



Other considerations for patients that are under 19 years old:

Child's Insurance Status	VFC Eligible?	VFC Eligibility Category
Has private health insurance plan with Medicaid as secondary insurance	Yes	Medicaid
Has health insurance covering all vaccines, but has not yet met plan's deductible or paid for other services received at visit AND has Medicaid as secondary insurance	Yes	Medicaid
Has private health insurance that covers all vaccinations and is American Indian/		AI/AN.
Alaskan Native (AI/AN)	Yes	However, provider should choose the eligibility category most cost-effective for the child and family.
		Medicaid or AI/AN.
Has Medicaid and is an American Indian/ Alaskan Native (AI/AN)	Yes	Provider should use Medicaid for the administration fee because this provides the least out-of-pocket expense for the family
		Underinsured.
Has insurance plan that does not cover all ACIP recommended vaccines		Child can only receive vaccines not covered by insurance plan.
		Insured.
Has health insurance covering all vaccines, but has not yet the plan's deductible or paid for other services received at visit		This applies even when the primary insurer would deny reimbursement for the cost of the vaccine and its administration because the plan's deductible has not been met.
Has health insurance covering all vaccines, but the plan has a fixed dollar limit or cap on amount it will cover	Depends	NO - Insured until the fixed dollar limit is met YES - Underinsured after the fixed dollar limit is reached

Indiana Department of Health-Immunization Division



VACCINE TRANSPORT

Required Vaccine Transportation Procedures

- If your office is moving locations and transporting vaccines, you MUST contact your IDOH Regional Quality Assurance Specialist. Relocating without notification may jeopardize VFC program enrollment status.
- Only appropriate packing materials (see following page) may be used to safely transport and/or temporarily store vaccines
 - If an acceptable portable refrigerator or freezer unit is used, unit preparation, packing and transport must follow the unit's guidelines (i.e. AcuTemp Coolers or Fridge-Freeze units).
 - Temporary storage (i.e. during unit cleaning; offsite clinics) is possible only when acceptable portable refrigerator or freezer units are used.

 Digital data loggers with continuous monitoring and recording capabilities <u>must be used</u> for all temperature monitoring during transport.

- Vaccines must be attended at all times during transport and/or temporary storage, including temperature documentation on the VFC Vaccine Emergency Transport Log.
 - If temperatures are not monitored during transport/temporary storage, the vaccine will be deemed wasted because there is no proof that temperatures remained within acceptable ranges, regardless of duration.
 - The responsible party will be held accountable to replace all wasted VFC doses per IDOH's Restitution policy.
- Packed vaccine may never be placed in the trunk of a vehicle.
- The receiving facility must immediately unpack and appropriately store vaccines into acceptable VFC vaccine storage units that are monitored using a DDL or other continuous monitoring system for which a MIN/MAX can be recorded once daily and point-in-time temperature recordings can be documented twice daily.

VARICELLA-CONTAINING VACCINES

The vaccine manufacturer does not recommend transporting varicella-containing vaccines (MMR/V,VAR). If these vaccines must be transported in an emergency situation:

- They should be packed and transported in a portable freezer unit that maintains the temperature between -58°F and +5°F (-50°C and -15°C).
- Dry ice is never allowed.
- If a portable freezer unit is not available and the vaccines cannot be packed in a unit that maintains +5°F or colder:
 - Varicella-containing vaccines may be transported and stored at refrigerator temperatures between 36°F and 46°F (2°C to 8°C).
 - o Refrigerated varicella-containing vaccines must be used within 72 hours.
 - Remaining refrigerated varicella-containing vaccine must be discarded if it is not used within 72 hours. This is considered a temperature excursion and must be documented on the Vaccine Incident Response Worksheet and a Vaccine Return form completed in VOMS. Refreezing varicella-containing vaccines that are stored at refrigerated temperatures is prohibited.



LogTag

Indiana Department of Health-Immunization Division



SUPPLIES FOR TRANSPORT

KEEP A COPY IN YOUR EMERGENCY TRANSPORT CONTAINER



Hard-sided coolers or Styrofoam™ vaccine shipping containers

- Coolers should be large enough for your location's typical supply of refrigerated vaccines.
- Can use original shipping boxes from manufacturers if available.
- · Do NOT use soft-sided collapsible coolers.



Conditioned frozen water bottles

- Use 16.9 oz. bottles for medium/large coolers or 8 oz. bottles for small coolers (enough for 2 layers inside cooler).
- Do NOT reuse coolant packs from original vaccine shipping container, as they increase risk of freezing vaccines.
- Freeze water bottles (can help regulate the temperature in your freezer).
- Before use, you must condition the frozen water bottles. Put them in a sink filled with several inches of cool or lukewarm water until you see a layer of water forming near the surface of bottle. The bottle is properly conditioned if ice block inside spins freely when rotated in your hand (this normally takes less than 5 minutes.



Insulating material — You will need two of each layer

- Insulating cushioning material Bubble wrap, packing foam, or Styrofoam[™] for a layer above and below the vaccines, at least 1 in thick. Make sure it covers the cardboard completely. Do NOT use packing peanuts or other loose material that might shift during transport.
- Corrugated cardboard Two pieces cut to fit interior dimensions of cooler(s) to be placed between insulating cushioning material and conditioned frozen water bottles.



Temperature monitoring device – Digital data logger (DDL) with buffered probe. Accuracy of +/-1°F (+/-0.5°C) with a current and valid certificate of calibration testing. Pre-chill buffered probe for at least 5 hours in refrigerator. Temperature monitoring device currently stored in refrigerator can be used, as long as there is a device to measure temperatures for any remaining vaccines.

Why do you need cardboard, bubble wrap, and conditioned frozen water bottles?

Conditioned frozen water bottles and corrugated cardboard used along with one inch of Insulating cushioning material such as bubble wrap keeps refrigerated vaccines at the right temperature and prevents them from freezing. Reusing vaccine coolant packs from original vaccine shipping containers can freeze and damage refrigerated vaccines.

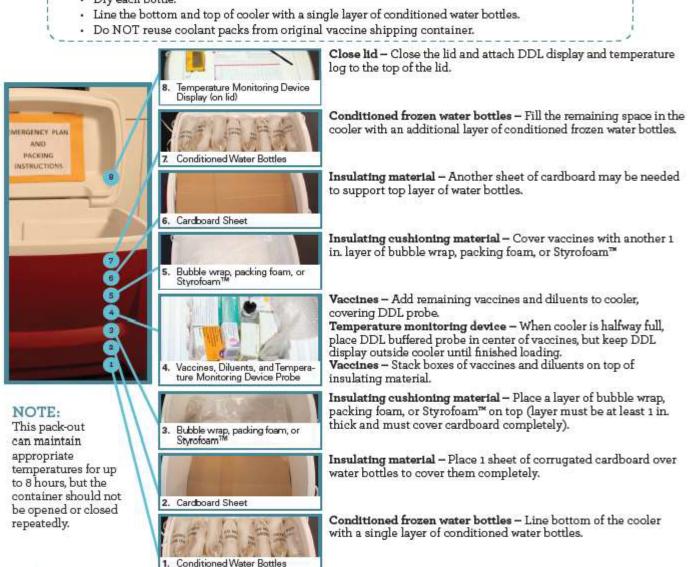
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PACKING THE VACCINE

Conditioning frozen water bottles (this normally takes less than 5 minutes)

- Put frozen water bottles in sink filled with several inches of cool or lukewarm water or under running tap water until you see a layer of water forming near surface of bottle.
- The bottle is properly conditioned if ice block inside spins freely when rotated in your hand.
- · If ice "sticks," put bottle back in water for another minute.
- Dry each bottle.



UNLOADING PACKED VACCINE

Arrive at Destination

- **Before opening cooler** Record time, temperature, and all needed information on the **VFC Vaccine Emergency Transport Log.**
- Storage- quickly transfer boxes of vaccines to appropriate storage units.
- **Troubleshooting-** If there was a temperature excursion during transport, label all affected vaccines DO NOT USE and store at appropriate temperatures.

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VACCINE TRANSFER PROCEDURES

Vaccine transfers involve the physical transfer of VFC vaccines and CHIRP inventory adjustments based on removal and receipt of VFC vaccines. Transfers are highly discouraged and should be rare events due to the possibilities of cold chain violations. Providers should use reminder recalls to avoid the transfer of vaccine whenever possible. Transfers of private-purchased vaccines are up to the discretion of the owning facility and are not subject to approval or guidance from IDOH.

As a standard practice, opened vaccine boxes should not be transferred. Partially used multi-dose vials cannot be transferred. It is the discretion of the IDOH Staff Representative to allow transfers, based on need and cost efficiency.

Transfer of vaccines should only occur for the following reasons:

- Unopened vaccine is three months (90 days) or less from expiration date, AND provider will not be able to use inventory prior to expiration date.
- An area outbreak has resulted in unexpected surge of walk-in patients.
- VFC clinic closure/disenrollment requiring vaccine redistribution to other VFC providers.
- Clinics needing to offload and transfer vaccines due to seasonal closure (such as a School Health Clinic).

If your office is moving locations and transferring vaccine, you MUST contact your IDOH Staff Representative. Relocating without notification may jeopardize VFC program enrollment status.

All VFC providers MUST obtain prior approval before transferring any VFC vaccine to another VFC provider*.

- Contact your IDOH Staff Representative to issue a transfer request. Please allow up to ten (10) business days for transfer approvals. All vaccine transfers are at the discretion of IDOH staff.
- All VFC vaccine transfers MUST follow IDOH vaccine transport procedures, including reporting of the transfer in VOMS.

^{*}Emergency vaccine transport to your backup storage facility is NOT considered a vaccine transfer and does not require advanced approval.

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OFFSITE CLINICS: VACCINE TRANSPORT, VACCINE USE, AND TEMPERATURE MONITORING PROCEDURES

Effective January 1, 2018, providers can no longer transport vaccines to/from offsite clinics in hard-side coolers or coolers available at general merchandise stores. All offsite clinics coordinated by VFC providers that plan on using VFC vaccine for patients must obtain approval from the IDOH Immunization Program at least two weeks prior to the event.

Offsite Clinic Requirements:

- 1. All clinics must use portable vaccine refrigerators/freezers or qualified pack-out units.
- 2. Digital data loggers with a buffered probe and a current and valid Certificate of Calibration Testing must be placed directly with the vaccines and used to monitor vaccine temperature during transport.
- 3. If vaccines are transferred to a permanent storage unit at the location of the offsite clinic, the storage unit must meet the minimum storage requirements for storage of VFC vaccines and the unit must have been monitored prior to the clinic day with a digital data logger.
- 4. Vaccines must be monitored during the clinic using a digital data logger <u>at least once an hour</u> and documented on the **Offsite Clinic VFC Vaccine Temperature Monitoring Log**.
- 5. Within the 24 hours following completion of the off-site clinic and return of all vaccines to the permanent storage unit, the data logger must be downloaded and the report must be reviewed and sent to the respective Ordering and Accountability Specialist.

Use of Multi-dose vials and Diluent at Offsite Clinics

- When a multi-dose vial is used, Food and Drug Administration (FDA) regulations require that it be used only by
 the provider's office where it was first opened. A partially used vial may be transported to or from off-site
 clinics operated by the same provider as long as the cold chain is properly maintained. However, such a vial
 may not be transferred to another provider or transported across state lines.
- Diluent should travel with its corresponding vaccine to ensure that there are always equal numbers of vaccine vials and diluent vials for reconstitution.
- Diluent should be transported according to its manufacturer guidelines and should not come into direct contact with conditioned water bottles because of the potential for freezing.

Varicella-Containing Vaccines Transport to Offsite Clinics

- CDC strongly discourages transport of varicella-containing vaccines to off-site clinics, because varicella-containing vaccines (VAR, Varivax; MMRV, ProQuad; ZOS, Zostavax) are sensitive to temperature excursions.
- Providers who choose to transport these vaccines to an offsite clinic must follow the appropriate procedures:
 - Transport in a portable freezer unit that maintains temperatures between -58°F and +5°F (-50°C and -15°C)
 - o The use of **dry ice is not allowed**, even for temporary storage.
 - Discard reconstituted vaccine if not used within 30 minutes.

EXCEPTION- REQUIRES APPROVAL

- Varicella-containing vaccines may be transported and stored at refrigerator temperatures at offsite clinics, between 36°F and 46°F (2°C to 8°C). This vaccine can be refrigerated for up to 72 continuous hours prior to reconstitution at the offsite clinic. IDOH must approve the number of anticipated doses for use.
- Varicella-containing vaccine stored at refrigerator temperatures must be discarded if it is not used within 72 hours. This is considered a temperature excursion and must be documented on the Vaccine Incident Response Worksheet and a Vaccine Return form completed in VOMS. Refreezing varicella-containing vaccines that are stored at refrigerated temperatures is prohibited

Indiana Department of Health-Immunization Division



Immunization Division Contact List

VFC Enrollment and Operations	
	Phone: (317) 416-5929
Contact for any analytic constitution of any artists	Fax: (317) 232-1913
Contact for general VFC enrollment questions.	Email: Enrollments@health.in.gov
	Available: Monday- Friday 8:00am to 4:00pm ET
VFC Quality Assurance Team	
Contact for VFC vaccine storage and handling issues,	Phone: (800) 701-0704
temperature excursions, VFC compliance questions or concerns,	Fax: (317) 233-3719
and training webinars, to report a facility change, or update the	Email: Immunize@health.in.gov
Primary and Back-up VFC Contacts	Available: Monday- Friday 8:00am to 4:00pm ET
Vaccine Ordering Management System (VOMS)	
Contact for vaccine ordering, inventory, reconciliation, returns,	Phone: (800) 701-0704
supply issues, VOMS training and VOMS user permissions.	Email: Immunize@health.in.gov
	Available: Monday- Friday 8:00am to 4:00pm ET
CHIRP Help Desk	
Control for a control CHIRD and the control of all the control of	Phone: (888) 227-4439
Contact for general CHIRP assistance, register a facility, and add or inactivate users.	Email: CHIRP@health.in.gov
of illactivate users.	Available: Monday- Friday 8:30am to 5:00pm ET
Immunization Division Document Center	Documents and forms can be found at the IDOH
	website at:
	https://www.in.gov/health/immunization/

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EMERGENCY VACCINE MANAGEMENT ACTION PLANS

Use the following guidance for emergency situations such as power outages, natural disasters, and equipment failure. This guidance also applies for any event that results in vaccine storage outside of recommended temperature range. This includes <u>responding to temperature excursions identified</u> <u>during DDL report reviews</u> and those <u>caused by human error (i.e. unit unplugged, unit door left open).</u>

USEFUL EMERGENCY NUMBERS

Service	Name	Phone #	E-mail
IDOH Staff Representative			
IDOH Immunization program		800-701-0704	vaccine@health.in.gov
Utility Company			
Building Maintenance			
Building Alarm Company			
Refrigerator/Freezer Alarm Company			
Refrigerator/Freezer Repair			
Point of Contact for Vaccine Transport			

EMERGENCY CONTACTS- In an emergency, contact the following people in the order listed:

Role/Responsibility	First & Last Name	Phone #	E-mail Address
1.			
2.			
3.			
4.			

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BACK-UP POWER SOURCES AND ALTERNATIVE LOCATIONS

Does the clinic have a generator?					
If so, (a) where is it locate (b) how often does it receitesting/maintenance?					
If your clinic does not have a generator, and/or your vaccine storage unit fails, it might be necessary to transport vaccines to an alternate storage location either within your building or offsite (e.g., a local hospital or another VFC provider).					
Identify an alternate loca meet VFC Program requi power outages affect you	rements. I	Note: <i>At least o</i>	_	=	_
Alternate Vaccine Storage Unit/Site	Physic	cal Address	Contact Person	Phone #	E-mail Address
Location of emergency p	oacking su	pplies			

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CHECKLIST: BEFORE AN EMERGENCY

Proper preparation for emergency situations is essential for protecting the viability of vaccines. Use the following checklist to help ensure practices are ready for planned or unexpected situations that might impact vaccines.

Step	Description
1.	Maintain current emergency contact information for key practice staff.
2.	Maintain current contact information for alternate vaccine storage location(s), including the facility name, address, and telephone number.
3.	Be familiar with backup power sources for your office (e.g. back-up generator).
4.	Stock vaccine packing and transport supplies, including back-up digital data loggers.
5.	Review (at least annually) the steps key practice staff must take to protect vaccines during all emergency situations, including how to pack and transport vaccines.

CHECKLIST: SCHEDULED POWER OUTAGE

Step	Description
1.	Notify the emergency contacts listed in this emergency plan and the IDOH staff representative.
2.	Contact an alternate storage facility to verify they can accept the vaccines and make a plan for transport. Update all contacts of proposed plan.
3.	Pack and transport vaccine per IDOH procedures.
4.	Once power has been restored, verify storage units are functioning properly and that temperatures have stabilized within acceptable temperature ranges.
5.	Notify and update emergency contacts.
6.	Follow the same vaccine transport procedures and return vaccine to its original storage unit.

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DURING AN EMERGENCY CHECKLISTS

Select the "During an Emergency" checklist that best applies to your emergency.

Power Outage/Natural Disaster

Follow these instructions during vaccine-related emergencies in compliance with VFC Program requirements and best practices.

Step	Description				
1.	Keep calm and do not open the unit.				
2.	Place a "DO NOT OPEN" sign on vaccine storage unit(s) and	d leave door(s) shut to conserve cold air mass.			
3.	Notify the emergency contacts listed in this emergency pla	ın.			
4.	Note the time the outage started and storage unit temper monitoring temperatures, even if they go out of range.	atures at the time of power loss; continue			
5.	Assess the situation to determine the cause of the power restore power.	Assess the situation to determine the cause of the power failure and estimate the time it will take to			
6.		 f the outage is short-term and/or will be tified in a less than 2 hours: Keep units closed and do not allow staff to use any vaccine. Do not transport or discard vaccine. 			
7.	Monitor vaccine storage unit temperatures until power is	restored.			
8.	Once power has been restored, follow all steps listed in After an Emergency.				

Human Error (Unit Unplugged/Unit Door Left Open)

Step	Description
1.	Upon hearing the alarm, identify and rectify the issue immediately (i.e. plug the unit in, shut the door).
2.	Keep calm and do not open the unit.
3.	Place a "DO NOT OPEN" sign on vaccine storage unit(s) and leave door(s) shut.
3.	Notify the emergency contacts listed in this emergency plan.
4.	Note the time the alarm started and storage unit temperatures at the time of alarm.
5.	Monitor the temperatures until they have stabilized to acceptable ranges.
6.	Follow the After an Emergency Step 6 if vaccines were exposed to out-of-range temperatures.
7.	Educate staff based on the identified human error issue (i.e. never unplug a vaccine storage unit).

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AFTER AN EMERGENCY CHECKLISTS

Power Outage/Natural Disaster/Human Error

Please follow these step-by-step instructions after vaccine related emergencies.

Step Description

- 1. Verify storage units are functioning properly.
- 2. Note the time that power was restored and/or temperatures stabilized within range, and document the storage unit temperatures (CURRENT, MIN and MAX) at that time.
- Once vaccine storage unit temperatures have stabilized, notify the emergency contacts.
- 4. If vaccines were transported due to an emergency situation:
 - A. Follow the same transportation procedures and transfer vaccine back to its original storage unit (Refer to the Vaccine Transport section for instructions).
- 5. If vaccines were maintained at required temperatures:
 - A. Remove the "DO NOT OPEN" sign from storage unit(s).
 - B. Notify staff that vaccines may be used.
- 6. If vaccines were exposed to out-of-range temperatures:
 - A. Label affected vaccines "Do Not Use" and keep them in the storage unit in acceptable temperatures- DO NOT remove them from the unit or discard the vaccine.
 - B. Document and report the excursion to your IDOH Staff Representative.
 - C. If vaccines are deemed spoiled by the manufacturer:
 - o Adjust inventory and submit return in VOMS.
 - o Place a vaccine order in VOMS .
 - Fill out the Temperature Excursion Reporting form and send it to your Ordering and Accountability Specialist.

When calling the manufacturer, you must speak to a person and obtain a case number for your temperature excursion. If the manufacturer is closed, keep the vaccines labeled, "Do Not Use," until you can get a case number.

Responding to an identified temperature excursion from DDL report review

Step Description

- 1. Keep calm and do not open the unit that was affected.
- 2. Place a "DO NOT OPEN" sign on vaccine storage unit(s) and leave door(s) shut.
- 3. Notify the emergency contacts listed in this emergency plan.
- 4. Note the time the excursion started and ended as well as the minimum and maximum temperatures reached during the excursion event.
- 5. Complete the Temperature Excursion Reporting Form and send it to your Ordering and Accountability Specialist

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- 6. A. If the storage units were the contributing factor to the excursion and need to be repaired, AND the vaccine is still viable:
 - 1. Contact an alternate storage facility to verify they can accept the vaccines.
 - 2. Refer to the Vaccine Transport section for instructions to safely relocate vaccines.
 - 3. Follow the After an Emergency: Storage Unit Repair/Replacement plan.

B. If the storage units were the contributing factor to the excursion and need to be repaired, AND the vaccine is **NOT** viable follow the **After an Emergency Step 6C.**

Vaccine cannot be stored in a combination unit when moving or changing units. Vaccine should be taken to a backup location and stored until 5 days of stable temperatures can be obtained.

Storage Unit Repair/Replacement

Step Description

- 1. Facilitate all storage unit repairs and or replacements. Involve your IDOH Staff Representative in any unit purchase to ensure it meets requirements.
- Once repairs are completed, or a new unit is installed, verify the unit is functioning properly.
 Record date of repair and update/add any unit information on the <u>Vaccine Storage Unit list</u> contained within this plan.
- 3. Monitor unit temperatures using a DDL/continuous monitoring device and record all temperatures on the corresponding unit temperature log. Water bottles should be placed in the unit at this time. It may take 2 to 7 days to stabilize the temperature in a newly installed or repaired unit. Do not move vaccines back to the unit until stable temperatures have been reached and approval received from your IDOH Staff Representative.
- 4. Once you have obtained 5 days of temperature recordings, submit temperature logs to your IDOH Staff Representative for review. Await instruction for transport or to continue monitoring the unit.
- 5. When approval is received, follow the <u>Vaccine Transport</u> guidelines and return vaccine to its original (or replaced, if applicable) storage unit.
- 6. Continue daily routine temperature monitoring.

The Centers for Disease Control and Prevention (CDC) requires all Vaccines for Children (VFC) providers to develop, maintain, and implement a Vaccine Management Plan with detailed and up-to-date standard operating procedures for routine and emergency vaccine management. The Indiana Department of Health (IDOH) Immunizations Division has prepared this document as a template to assist your office with this requirement. Please refer to IDOH VFC Program Policies for supplemental information and guidance.

For access to the most recent policies, documents and changes, as well as, IDOH staff directory and maps, please go to https://www.in.gov/health/immunization/

Vaccine Storage Troubleshooting Record

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Use this form to document any unacceptable vaccine storage event, such as exposure of refrigerated vaccines to temperatures that are outside the manufacturers' recommended storage ranges. Send this form to your IDOH Ordering and Accountability Specialist.

Date & Time of Event If multiple, related events occurred, see Description of Event below.	Storage Unit Temperature at the time the problem was discovered		Room Temperature at the time the problem was discovered	Person Completing Report	
Date:	Temp when discovered:		Temp when discovered:	Name:	
Time:	Minimum temp:	Maximum temp:	Comment (optional):	Title:	Date:
Description of Event (If multiple, related events occurred, list each date, time, and length of time out of storage.) General description (i.e., what happened?) Estimated length of time between event and last documented reading of storage temperature in acceptable range (2° to 8°C [36° to 46°F] for refrigerator; -50° to -15°C [-58° to 5°F] for freezer) Inventory of affected vaccines, including (1) lot #s and (2) whether purchased with public (for example, VFC) or private funds (Use separate sheet if needed, but maintain the inventory with this troubleshooting record.) At the time of the event, what else was in the storage unit? For example, were there water bottles in the refrigerator and/or frozen coolant packs in the freezer? Prior to this event, have there been any storage problems with this unit and/or with the affected vaccine? Include any other information you feel might be relevant to understanding the event.					
Action Taken (Document thoroughly. This information is critical to determining whether the vaccine might still be viable!)					ıfacturers
 When were the affected vaccines placed in proper storage conditions? (Note: Do not discard the vaccine. Store exposed vaccine in proper conditions and label it "do not use" until after the manufacturers determine viability and you can obtain a case number). Who was contacted regarding the incident? (For example, supervisor, state/local health department, manufacturer—list all.) IMPORTANT: What steps did you take to prevent a similar issue from happening in the future? 					nacturers
Results What happened to the vaccine? Did the manufacturers tell you to use or waste the vaccine? Please record the Case Number here. Send this form to your Ordering and Accountability Specialist.					

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COVID-19 Vaccine Addendum

COVID-19 Vaccination Provider Requirements

All COVID-19 vaccination providers participating in the <u>U.S. COVID-19 Vaccination Program</u> are required to sign a COVID-19 Vaccination Program Provider Agreement to receive delivery of any COVID-19 vaccine from CDC's distributor or a COVID-19 vaccine manufacturer. The agreement must be completed by all public and private providers, provider organizations, and government-affiliated federal, state, territorial, and local providers.

As part of the agreement, providers are required to:

- Store and handle COVID-19 vaccines under proper conditions, including maintaining cold chain conditions and chain of custody at all times in accordance with an EUA or vaccine package insert, manufacturer guidance, and CDC guidance in this toolkit.
- <u>Monitor storage unit temperatures</u> at all times, using equipment and practices that comply with guidance in this toolkit.
- Comply with immunization program guidance for handling <u>temperature excursions</u>.
- Monitor and comply with COVID-19 vaccine expiration dates.
- Preserve all records related to COVID-19 vaccine management for a minimum of three years.
- Comply with federal instructions and timelines for disposing of COVID-19 vaccine and diluent, including unused doses.

COVID-19 Vaccine Transport

As part of the COVID-19 Vaccination Program, a minimum order size of COVID-19 vaccine, diluent (if applicable), and ancillary supplies will be shipped directly to enrolled COVID-19 vaccination providers. In most instances, vaccine will be delivered directly to the facility where it will be administered to maintain the vaccine cold chain. However, there may be circumstances where COVID-19 vaccine needs to be redistributed or transported. This can occur if:

- A CDC Supplemental COVID-19 Vaccine Redistribution Agreement is in place for vaccine to be redistributed beyond the identified primary CDC ship-to site (i.e., for larger organizations whose vaccine is shipped to a central depot and requires redistribution to additional clinic locations).
- A jurisdiction immunization program or commercial partner needs to redistribute vaccine within the jurisdiction based on need or to keep vaccine from being wasted.
- Vaccine needs to be transported for satellite, temporary, or off-site clinics, including programs at long-term care facilities, or for administration to patients such as those in a home healthcare program.

In these instances, appropriate precautions must be taken to protect the vaccine. Vaccine must only be transported using appropriate packing materials that provide maximum protection. Follow specific jurisdiction and federal direction for transporting COVID-19 vaccine products.

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COVID-19 Vaccine Transport Continued

The chart below shows general transport recommendations to maintain the vaccine cold chain in two situations: emergency transport and transport for use at off-site clinics or satellite facilities or for relocation of stock. Recommendations vary based on the situation. Some COVID-19 vaccine products may have specific transport guidance to ensure the cold chain is maintained and vaccine is protected.

General Transport System Recommendations	Emergency Transport	Transport for Off-Site Clinic, Satellite Facility, or Relocation of Stock	
Portable Vaccine Refrigerator, Freezer, or Ultra-cold Freezer	Yes	Yes	
Qualified Container and Packout	Yes	Yes	
Conditioned Water Bottle Transport System	Yes	No	
Manufacturer's Original Shipping Container	Yes (last resort only)	No*	
Food/ Beverage Coolers	No	No	

^{*}The original shipping container for ultra-cold COVID-19 vaccine can be used for transport.

COVID-19 Vaccine Emergency Storage and Handling

Emergencies such as equipment failures, power outages, severe weather conditions, or natural disasters usually happen without warning and may compromise storage conditions. Because of the limited supply of COVID-19 vaccine, especially during early U.S. vaccination efforts, it is critical that COVID-19 vaccination providers have plans in place for emergency situations. Some key issues to remember include:

- Vaccines may remain inside a nonfunctioning unit as long as appropriate temperatures are maintained.
 Monitor your DDL to determine when additional action should be taken.
- Having an on-site generator(s) prevents the need to transport vaccines to an alternative storage facility during a power outage.
- Emergency situations can arise outside of normal business hours, so your office staff as well your facility's building manager and/or security staff, if appropriate, must understand how to implement your emergency operation plans or access your facility if necessary.
- Ensure your facility has the resources on hand to safely pack vaccines for transport during emergencies.

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Signature Page:

I have read through the current Vaccine Management Plan. I fully understand this practice's responsibilities regarding publicly funded vaccines.

Employee Signature	Title/Position	Date

Vaccine Emergency Management Plan

Post on the outside of each refrigerator and freezer

Practice Name:	PIN:	
Address and Office		
Phone Number:		
Revision Date:		
Primary Coordinator:		
Phone Number:		
Work and Home/Cell Phone)		
Back-Up Coordinator:		
Phone Number:		
Work and Home/Cell Phone)		
Person with 24-hour Access:		
Phone Number:		
(Work and Home/Cell Phone)		

Ensure primary and back-up contacts are on a notification list for practice-based power outages.

efore transporting, call back-up lo	cation to ensure their generator is working.
1. Location and Contact Name:	
Phone Number:	
2. Location and Contact Name:	
Phone Number:	
accine should only be transported	or shipped in an emergency and according to the VFC Vaccine Storage and Handling Toolkit o/admin/storage/toolkit/storage-handling-toolkit.pdf).
mergency Contacts	
Refrigeration Company:	
Phone Number:	
Utility Company:	
Phone Number:	
Other—Describe:	
Phone Number:	
Other Resources	
Regional Quality Assurance Specialist:	
Phone Number and E-mail:	
Vaccine Ordering and Accountability Specialist	
Phone Number and E-mail:	

For a Power Outage: If you do not have a back up generator, identify one location with a generator (hospital, 24-hour store, etc.)

If temperature excursions are noted, please notify your IDOH Staff Representative or call the Immunization Division.

Immunization Toll Free:

1-800-701-0704



