Indicator Reference Sheets

This section describes the primary key indicators. Note that the selection of indicators to include in a dashboard is context specific. No indicator is to be considered more useful or of higher value than any other.

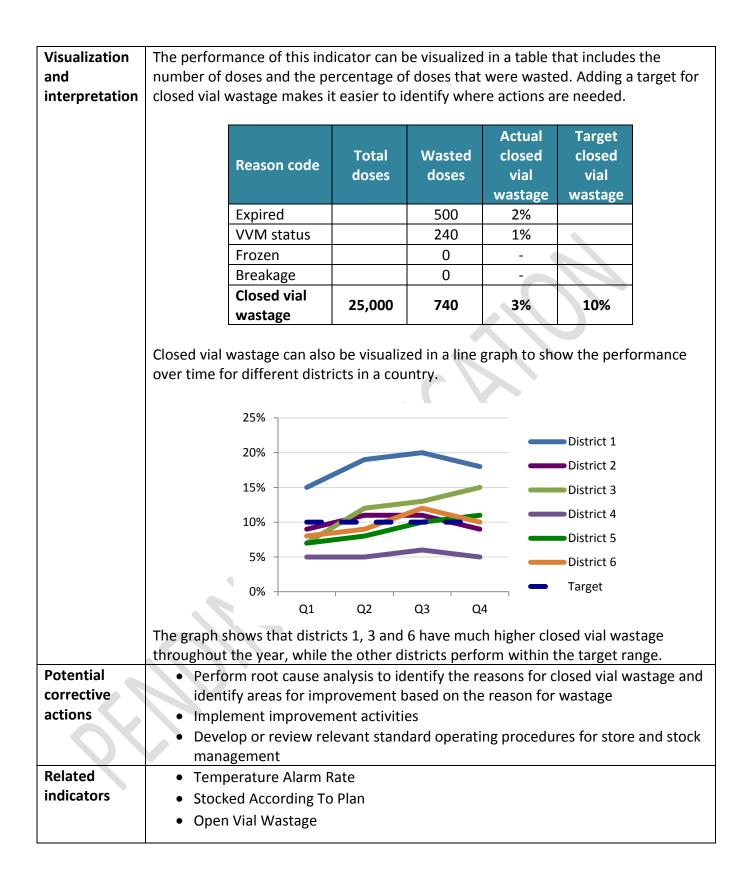
A short description of the type of information found in each sheet follows:

- The **name of the indicator** is at the top, along with a **description** of the indicator and the **purpose** of measuring it. The purpose section includes the questions a manager might ask that the indicator could answer.
- The **performance objective** refers to the strategic objective with which the indicator is most closely associated (vaccine availability, vaccine potency and supply chain efficiency), while the **domain** is the supply chain component to which the indicator belongs.
- **Full indicator name(s)** reflects the specific ways the indicator can be calculated depending on the managers who will use the dashboard. Most of the indicators can be calculated differently at each supply chain level: for example, by aggregation, by health facility or by district.
- **Dashboard use level** refers to the level(s) of the supply chain where the dashboard is recommended for use.
- **Preconditions** lists any conditions (e.g., policies, data availability) that might need to be in place in order to implement and use the indicator, and **system design** specifies the type of system (e.g., push, pull systems) where the indicator is relevant.
- The data needed, data sources and data collection method sections provide details of those topics, while the calculation section includes formulas and examples to better illustrate calculation of the indicator, and the visualization and interpretation section includes examples relevant to different supply chain levels.
- **Potential corrective actions** might be triggered by an extraordinary performance value; they are divided into 'operational' and 'strategic' management actions where appropriate. Operational actions involve the routine management of the supply chain ensuring that products are in stock, that temperatures are maintained and that the system is performing as expected. They often focus on how to address a particular problem directly. Strategic management actions, on the other hand, are typically more long-range, involving high-level decisions about system design, planning and procurement, and often focusing on how to prevent a particular problem from recurring.
- **Related indicators** are added to provide guidance on expanding the dashboard or determining which diagnostic indicators would be required for root cause analysis. They also show which of the other primary key indicators are specifically related to the indicator in question.

Name	Closed Vial Wastage
Description	Percentage of the total number of closed vial vaccine doses managed by a store or health facility during a particular period that are spoiled because of expiry, heat exposure, freezing, breakage, loss of the accompanying diluent or discard of unopened vials at the end of an outreach session. Wastage at the point of administration, because of incomplete use of the contents of a multi-dose vial, is referred to as open vial wastage and is not included in closed vial wastage ¹ .
Purpose	The indicator is used to measure potential avoidable wastage during transportation and storage. Wastage is related to the performance of vaccine ordering, distribution and store management. It can indicate excessive ordering practices that are not well- aligned to actual consumption rates, vaccine exposure to heat or freezing temperatures, breakage and mishandling of inventory.
	 This indicator can help answer questions such as: How many extra vaccines should be procured beyond those estimated to be administered? Do the quantities of vaccine ordered at particular facilities routinely exceed actual usage? What is the approximate financial value of closed vial wasted vaccine? Is wastage similar between facilities and between districts? Is targeted reinforcement of standard operating procedures and vaccine management principles needed?
Performance	Availability
objective	Potency Efficiency
Domain	Stock management
Full indicator	 Closed vial wastage rate per facility
name(s)	 Average closed vial wastage rate
	 Closed vial wastage rate per district/administrative level
Dashboard	This indicator is recommended in dashboards used by sub-national and national
use level	managers and by store managers at all levels.
Preconditions	A system for recording closed vial wastage, optionally with reason codes, needs to be
System design	in place. Relevant in all types of logistics systems.
-	
Data needed	 Number of discarded (wasted) doses reported by vaccine and preferably by reason code Number of doses under management during a certain period, defined as the starting balance plus all of the doses received during that period
Data sources	 Vaccine stock ledgers/cards Vaccine orders Batch management to track vaccine vial monitor (VVM) status and expiry dates Logistics management information system (LMIS)

¹ Further information on wastage can be found at <<u>http://apps.who.int/iris/bitstream/10665/68463/1/WHO_VB_03.18.Rev.1_eng.pdf</u>>.

	Wastage reporting tools
Calculation	Closed vial wastage = $\frac{\text{number of doses discarded during reporting period}}{\text{doses under management during the same period}} \times 100$
	Doses under management is defined as the opening balance plus all doses that were received during the period. Issued doses should not be subtracted.
	 Closed vial wastage should include vials wasted due to: Expiry, which may indicate ordering practices that are not aligned to actual consumption rates, failure to respect first expiry first out (FEFO) policies, a supply design that moves too slowly (i.e., it takes too long for a vaccine to go through the chain to the point of administration) or poor organization in a vaccine store such that an older lot or batch can be overlooked. VVM status 3 or 4 (at or beyond the discard point) before the vaccine's expiry date has been reached, which may indicate poor cold chain quality or breaches in the cold chain. Freezing, which is an indication of poorly functioning cold chain equipment or poor adherence to standard operating procedures during storage or transportation.
	• Breakage, either of the vials or accompanying diluent. Inclusion of reason codes in reporting of closed vial wastage allows additional
	precision and more thorough investigation of root causes.
	Example In a regional store, 500 doses of pentavalent vaccines expired during the year and 240 doses were wasted due to VVM status 3 or 4, bringing the total for the period to 740 doses.
	If the beginning balance of pentavalent vaccines in that same store was 5,000 doses, and four shipments of 5,000 doses were received during the year, then the total number of doses under management during the year was 25,000 doses.
0	Closed vial wastage (pentavalent) = $\frac{740 \text{ doses}}{25.000 \text{ doses}} \times 100 = 3\%$
	When calculating by reason code, the overall closed vial wastage is divided into:
	Closed vial wastage due to expiry (pentavalent) = $\frac{500 \text{ doses}}{25.000 \text{ doses}} \times 100 = 2\%$
	Closed vial wastage due to VVM status 3 and 4 (pentavalent) = $\frac{240 \text{ doses}}{25.000 \text{ doses}} \times 100 = 1\%$



Name	Forecasted Demand Ratio
Definition	Ratio of actual consumption of a given product during a particular period compared to the consumption forecasted for the same period. Consumption includes administered and wasted doses.
Purpose	Used to validate and improve forecasting practices and assumptions (e.g., target population, coverage, wastage) in order to increase forecasting accuracy.
	 The indicator helps to answer questions such as: Is consumption in a health facility, administrative unit or country as expected? Is there a need to plan for additional stock to avoid stock-outs? Is closed vial wastage likely due to lower usage than expected? Is there a need to review the forecasting assumptions (e.g., target population, coverage)? Is there a need to revise minimum and maximum stock levels?
Performance objective	Availability
Domain	Demand planning
Full indicator	Health facility forecasted demand ratio
name(s)	Average forecasted demand ratio for sub-national level
	 % of health facilities with forecasted demand ratio in a set interval
Dashboard	This indicator is recommended in dashboards used by sub-national and national
use level	managers.
Preconditions	Consumption (i.e., administered and wasted doses) data is necessary to calculate the
	indicator, so a system to collect actual consumption data is necessary.
System	Relevant in all types of supply chain systems.
design	
Data needed	 Forecasted demand/usage by product Actual consumption by product (opening balance + receipts – closing balance of product)
Data sources	 Logistics management information system (LMIS)
	Monthly immunization reports
	Micro plans
	Stock ledgers/cards
Calculation	Forecasted demand ratio = (doses consumed per product in a period/doses forecasted per product for the same period)
	It is important that the doses consumed and the doses forecasted apply to the same period. The longer the period, the more accurate the forecasted demand ratio. A rolling year, half year or quarter are recommended, but the length of the period might depend on the reliable data available and the staff's ability to calculate indicator performance for a long period. Interpreting the ratio:

		1: actual consumption (through s less than the forecasted consumption for a
	given period.	4
	 Forecasted demand ratio above a administration and wastage) was 	s more than the forecasted consumption for
	a given period.	
	 A forecasted demand ratio close 	to 1 implies that the forecasted
	consumption matched well with	-
	Average forecasted demand ratio = $(\sum h)$ ratios)/(total # health facilities)	ealth facility forecasted demand
	-	he percentage of facilities with a forecasted or example, within the range of 0.7 to 1.3).
	Evample	
	<i>Example</i> In a health facility, the quarterly forecas vials was 45 vials, whereas the actual co quarter was 35 vials.	ted usage of yellow fever vaccine in 10-dose nsumption of this vaccine in the same
	Forecasted demand ratio for yellow feve	er vaccine = 350/450 = 0.78
	The forecasted demand ratio shows that lower than forecasted (forecasted dema	t the health facility's actual consumption was nd ratio < 1).
		orecasted usage of yellow fever vaccine in I consumption of this vaccine in the same
	Forecasted demand ratio for yellow feve	er vaccine = 450/400 = 1.13
	For this health facility, actual consumpti	on was higher than the expected
		stock had to be used (forecasted demand
6	District A is preparing its district report, reported their forecasted demand ratio	6
	Health facility	Forecasted demand ratio
	Health facility 1	0.78
	Health facility 2	1.13
	Health facility 3	1.50
	Health facility 4	1.25
	Health facility 5	0.85
	Health facility 6	0.93
	Health facility 7	0.98

	Average forecasted demand ratio (district A) = (0.78 + 1.13 + 1.50 + 1.25 + 0.85 +		
	0.93 + 0.98)/7 = 1.06		
	The average forecasted demand ratio shows that the overall district consumption is close to the consumption forecasted.		
	Another way to report the aggregated forecasted demand ratio is to calculate the percentage of health facilities with usage within set limits. In this example, a +/– 20% ratio is used. This method of calculation more clearly shows how many health facilities are consuming more or less than expected.		
	% of health facilities with forecasted demand ratio between 0.8 and 1.2 = $4/7 \times 100 = 57\%$		
	57% of the health facilities in District A have consumption within the set target interval. The remaining facilities have either higher or lower usage than expected.		
Visualization	The forecasted demand ratio can be visualized in a bar chart. Values above 1 indicate		
and	consumption above forecasted demand quantities, while a ratio below 1 indicates		
interpretation			
	l identify the health facilities where actual usage differs from forecasted usage		
	identify the health facilities where actual usage differs from forecasted usage.		
	identify the health facilities where actual usage differs from forecasted usage. Forecasted demand ratio for yellow fever (10-dose vial)		
	Forecasted demand ratio for yellow fever (10-dose		
	Forecasted demand ratio for yellow fever (10-dose vial)		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.95		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.75 0.5 0.5		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.95		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.5 0 HF1 HF2 HF3 HF4		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.95 0.5 0 0 1 0.95		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.75 0.5 0 HF1 HF2 HF3 HF4 Low target High target Forecasted demand ratio		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1 0.5 0 HF1 HF2 HF3 HF4		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1.5 0.5 0.75 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0.		
	Forecasted demand ratio for yellow fever (10-dose vial) 1.5 1.5 1.5 0.5 0.5 0.5 0.5 0.5 0.5 0.5 0		

	2 — 1.5 — 1 — 0.5 — 0 —	Q1	Q2	Q3 District 3	LO-dose vial) Q4 District 4	
	the districts, bu further. A spatial displa example, the co facilities have a illustrative and District D	shows that the ut in District 1, t y is another wa olours indicate a forecasted der can be adapted District B Red for Gre wit	forecasted den here is a varia y to give an ov whether more mand ratio bet d to the contex d = District wit ecasted deman een = District w h forecasted d	mand ratio is fa tion over time t verview of distr or less than 80 tween 0.8 and 2 kt. h <80% of heal nd ratio betwee vith >80% of heal	hirly constant in t that could be inv ict performance. 0% of a district's 1.2. The targets t th facilities with en 0.8 and 1.2	vestigated . In this health
Potential corrective actions	 Review identify tolerand could be inaccura Revise of consister Revise r consister 	reasons for for ce level (e.g., sto e: inaccurate as ate on-time and ordering policies ently outside of minimum and mently too high of	methodology ecasted demain ock-out can lea sumptions (tan l in-full deliver s and practices the tolerance naximum stock r too low	and perform a nd ratios beyor ad to a forecast rget population ies, higher was s when the fore level or there is	root cause analy nd the establishe ratio < 1). Root , coverage and w tage than expect casted demand s a large imbalan precasted deman	d causes wastage), ted. ratio is nce
Related indicators	Full StoreClosed V	l According to P ck Availability Vial Wastage e and In-Full De				

Name	Full Stock Availability
Description	Percentage of storage points with full availability of all or a selected set of tracer vaccines and immunization supplies over a resupply period. Full availability is defined as no stock-out in the store or health facility at any point during the time period.
Purpose	 Measures the availability of immunization products. Availability of vaccines and immunization supplies is important to reach immunization programme targets. The following questions can be answered by monitoring the performance of this indicator: Are certain facilities frequently at risk of stock-outs? What is the full availability percentage by district or region? Does low availability in the national or resupply store affect availability at
	lower levels?Is full availability lower than expected in certain health facilities or regions?
Performance objective	Availability
Domain	Stock management
Full indicator	 % of health facilities with full availability
name(s)	% of districts with full availability
	% of districts with at least x% of facilities with full stock availability
Dashboard use level	This indicator is recommended in dashboards used by sub-national and national managers.
Preconditions	This indicator can be implemented in any context, as it requires only observation of zero stock balance during the resupply period.
System design	Relevant in all types of logistic systems.
Data needed	 Product stock-outs in stores and health facilities OR: closing balances at the end of the resupply period in stores and health facilities
Data sources	 Stock cards/ledgers Physical inventory/physical stock counts Stock-out reports from health facilities Logistics management information system (LMIS)
Data collection method	Where necessary, full availability can be determined for a basket of tracer indicator products representing the availability of immunization supplies.
Calculation	Full stock availability = resupply periods without stock-out of any (tracer) vaccine or immunization supplies
	At sub-national and national level, the indicator is aggregated as % of health facilities or % of districts with full stock availability. The calculation for a sub-national region is:
	% health facilities with full stock availability = (# health facilities with full availability of all (tracer) immunization products)/(total number of health facilities in sub-national region) x 100

Alternatively, for the national level, the aggregation can be based on the percentages of health facilities in a district exceeding a set threshold.

Districts with full availability of all (tracer) immunization products in more than x% of health facilities = (# districts with more than x% health facilities with full availability of all [tracer] immunization products in the last resupply period)/total # districts) x 100

The percentage of health facilities in the above calculation is set by the country to reflect the expected standards. When reporting the value of the indicator, the threshold value must be included.

Example

The table below shows health facility A's report to the district on stock availability for the country's tracer immunization products in the second quarter (Q2). Deliveries to the facility are monthly.

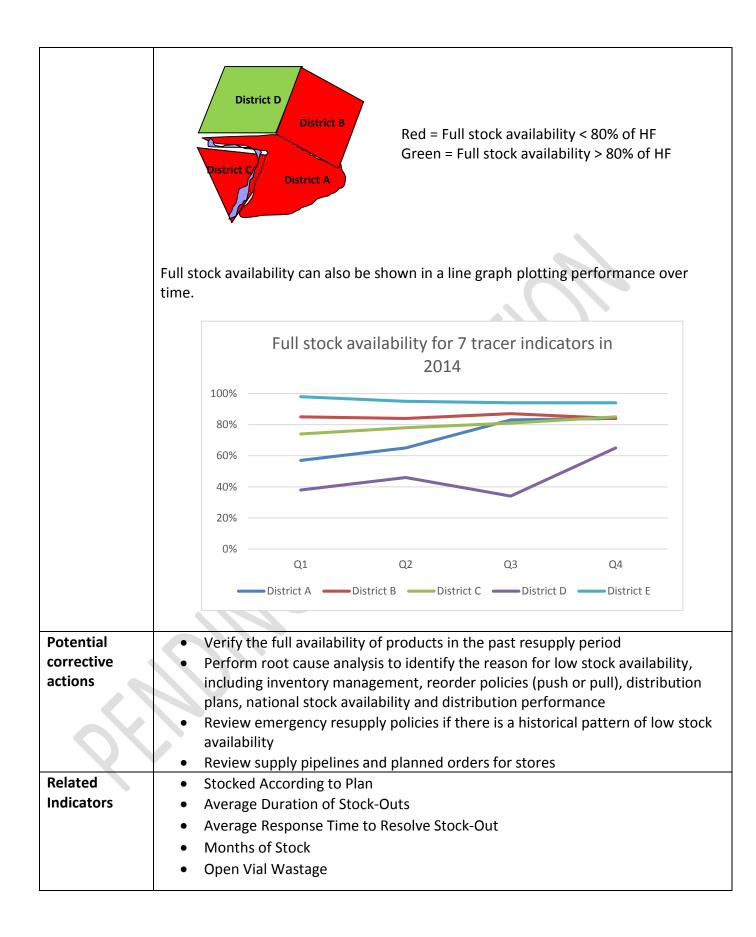
Tracer	Vaccines available			
immunization products	April	Мау	June	
BCG	YES	YES	YES	
PCV	YES	NO	NO	
Pentavalent	YES	NO	NO	
Rotavirus	YES	YES	YES	
Syringe 0.5 ml	YES	YES	YES	
Measles	YES	YES	YES	
Full availability?	YES	NO	NO	

According to the table, there was full availability of all tracer immunization supplies in health facility A in April, but in both of the other months, at least one vaccine was not fully available. Therefore, health facility A had full stock availability only in April.

When the full stock availability percentages for each district are received at the national level, the national stock availability can be calculated as a national average or as a percentage of districts above a set percentage of health facilities with full availability.

District	# of health facilities with full availability	Total # of health facilities	Q2
District A	6	15	40%
District B	10	16	63%
District C	15	21	71%
District D	10	12	83%
District E	18	19	95%

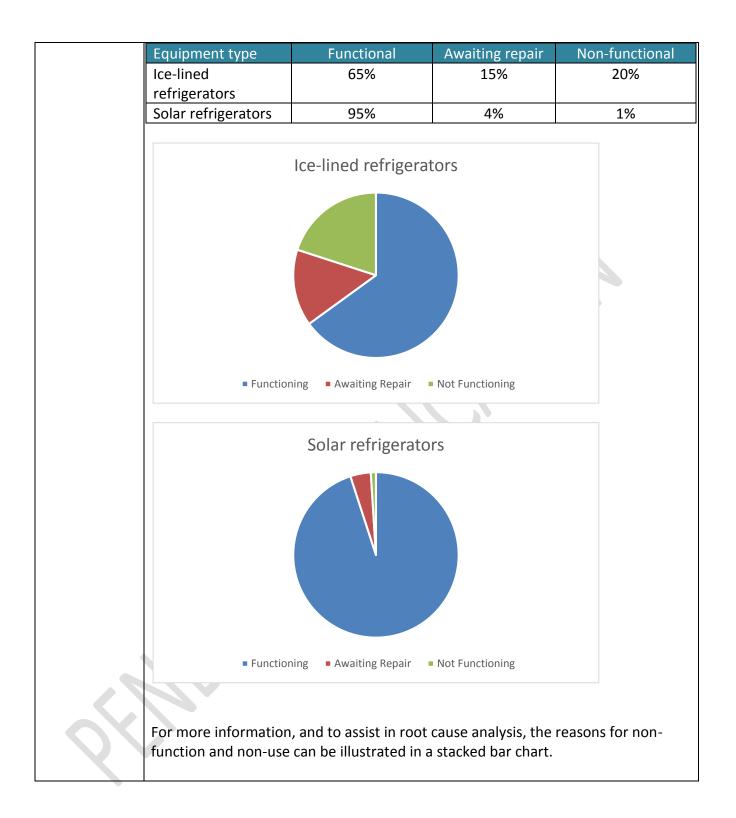
	District F	15	18	83%
	District G	9	11	82%
	District H	16	24	67%
	District I	16	21	76%
	District J	16	16	100%
	District K	15	18	83%
	National full stock availability	146	191	76%
Visualization	National full stock availa health facilities x 100 = 1 The country has set 80% availability. % districts with at least 8 with >80% health facilitie 100 = 55% A colour-coded table car	46/191 x 100 = 76% as the defined three 80% of health facilitie es with full stock ava	shold for health facili es with full stock ava iilability/total # distr	ities with full stock ilability = # districts icts) x 100 = 6/11 x
and				-
interpretation	threshold for green and red performance has to be set according to the context and the availability. Here, 80% was used as the threshold.			
		District	Q2	
		District A	40%	
		District B	63%	
		District C	71%	
		District D	83%	
		District E	95%	
		District F	83%	
		District G	82%	
		District H	67%	
	Another way to visually in through colour-coded sp again used as the perform health facilities (HF) have supplies, while in the gree	atial analysis. In the mance threshold. In e full availability of a	visualization of a reg the red districts, feven set of tracer vaccine	gion below, 80% was wer than 80% of es and immunization

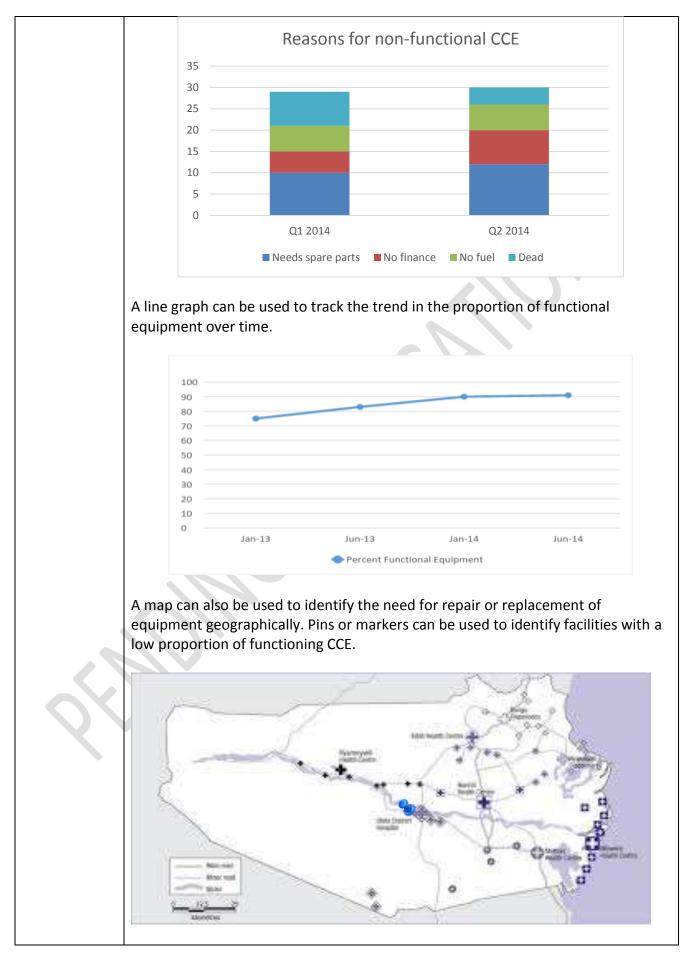


Name	Eurotional Status of Cold Chain Equipment
	Functional Status of Cold Chain Equipment
Description	Cold chain equipment functioning compares the proportion of cold chain equipment (CCE) operable for storing vaccines with the overall number of commissioned CCE devices in a particular area. CCE is defined as all refrigerators, freezers, passive storage devices, and walk-in cold rooms and freezer rooms designated for storing vaccines. CCE functioning can be measured at a point in
	time or over a particular period of time.
Purpose	Measures functional cold chain equipment to identify risk of inadequate cold storage for maintaining vaccine potency. Used for operational purposes, such as to ensure that vaccines are appropriately stored (e.g., to target deployment of repair or maintenance services), and for strategic purposes, such as to plan for replacement of equipment or parts.
	Over time, the trend in the proportion of functional equipment can be used to measure performance of in-house or contracted maintenance and repair services. If the proportion of functional equipment is disaggregated by reason for the non-function or by equipment type, the indicator can also be used to assess the performance of particular types or models of CCE in the field.
	Note that functional status of CCE does not include a provision regarding the temperature maintained by the equipment; other indicators (such as Temperature Alarm Rate) must be used to fully understand the cold chain management system.
	 The following questions can be answered by monitoring this indicator: Which CCE is in need of repair or maintenance and where is it located? In case of delivery of additional CCE, where is it most needed? What investment in new CCE is needed for the next few years? Do particular models or types of CCE perform more reliably or have a longer lifespan than others?
Performance objective	Potency
Domain	Cold chain management
Full indicator name(s)	 % of functional CCE % of health facilities or % of districts meeting a threshold for functional CCE (e.g., % of districts with at least 90% functional equipment)
Dashboard use level	This indicator is recommended in dashboards used by sub-national and national managers and all store managers.
Preconditions	The indicator requires an updated cold chain equipment inventory, a mechanism to ascertain whether equipment is functioning and a system to transmit the information to the level where cold chain equipment planning is undertaken. The transmission mechanism can be paper-based, electronic or communication-based (e.g., by telephone).
System design	Relevant in all types of logistics systems.

Data needed	 Number of CCE devices designated for storing vaccines in a particular geographical area Functional status of each CCE: functioning/awaiting repair/unserviceable Primary reason for not functioning or not in use: needs spare parts/no finance/no fuel/surplus/dead/not applicable Optional additional data: temperature of CCE Note: Precise definitions of the functional status and reasons for not functioning need to be standardized to allow comparison. For instance, CCE operating outside the normal range of temperature may be considered awaiting repair. Power sources, such as generators for backup power for walk-in cold facilities, may also be included in this indicator, as appropriate. Multiple reason codes may also be applied.
Data sources	Cold chain equipment inventories by location; for example, WHO Excel-
Data sources	 Cold Chain Equipment Inventories by location, for example, who Excel- based tool, Cold Chain Equipment Inventory (CCEI), Cold Chain Equipment Manager (CCEM) On-site assessment of equipment functioning Maintenance worksheet CCE distribution plan
Calculation	% CCE functioning = (# functioning CCE devices)/(total # CCE devices designated for
	use in reporting facilities) x 100 The indicator can be calculated either at a point in time or over a period of time. When calculated over a period of time, % CCE functioning needs to take into account how long the non-functional periods were: % CCE functioning = number of functional CCE unit-days/total number of CCE unit- days in a given reporting period x 100, where CCE unit-days are the total number of days in the reporting period multiplied by the number of CCE devices. Both the numerator and the denominator should be collected from the same geographic area, and decommissioned equipment should not be counted in either the numerator or denominator. Functionality of CCE is broadly meant to mean that the device is operable at a particular point in time for storing vaccine. Disaggregation of both the numerator and denominator by location and by type, manufacturer, model, energy source, PQS (performance, quality and safety) code or year of installation can add value in investigating root causes of CCE failures, in targeting maintenance and replacement, and in performance monitoring of equipment and of maintenance systems. Examples Consider a district with 50 facilities. The most recent CCE inventory indicates that the following equipment is available in the district, and a recent facility survey found the following number and percentage of devices functional:

		Туре	Total number	Number	Percentage				
				functioning	functioning				
		Ice-lined refrigerators	35	25	71%				
		Deep freezers	5	3	60%				
		Solar direct	15	14	0.2%				
		drive refrigerators	15	14	93%				
		onally, the district d in the above ta	had 3 decommiss ble.	sioned refrigerato	rs that were not				
	Overal	l:							
	(42 fur	nctional devices)/	(55 total devices)	x 100 = 76% funct	ioning				
				\sim					
			•		CE inventory and a				
	facility	survey found the	following equipm	ent functional:					
					Deveevters				
		Region	Total number	Number	Percentage				
		Α	100	functioning 95	functioning 95%				
		A	200	184	92%				
		C	150	149	99%				
		D	300	265	88%				
		Е	85	73	86%				
	Overall: (766 functional devices)/(835 total devices) x 100 = 92% functioning (3 regions with >90% CCE functioning)/(5 total regions) x 100 = 60% of regions with >90% CCE functioning								
Visualization			l, data from all fac	cilities or an aggre	gate across facilities	in			
and					art or sorted table ca				
interpretation	display	the proportion o	of equipment that	is functional, and	managers can then u	use			
			itor performance	of maintenance s	ystems and of particu	ula			
	types o	or models of CCE.				1			
	Fa	acility	Total number of	Number	Percentage				
			CCE	functioning	functioning				
		ealth centre A	2	2	100%				
		ealth centre B	3	2	67%	-			
		ealth centre C	1	0	0%	-			
		otal	6	4	67%	l			
	-	rformance of par or a pie chart.	ticular types of CC	E in a district can	also be visualized in	а			





	At the national level, data are further aggregated across sub-national administrative regions or as a single estimate of functionality nationwide. Pie charts and line graphs are also useful at the national level.
Potential	Verify that equipment is not functioning
corrective	
actions	 Determine the root cause of equipment dysfunction; solicit repair or replacement of non-functional equipment
	• Ensure that contingency plans are in place for all facilities, so that vaccines
	can be safely stored or transported elsewhere when one or more devices are non-functional
	 Perform routine maintenance of all CCE to prevent future breakdown
	 Use equipment status (including reasons) to inform future procurement decisions
	 Reallocate functional CCE equitably, as appropriate
	• The indicator can also be used in combination with other inputs, such as a
	cold chain inventory, to estimate the total volume of cold chain space
	available and is useful in assessing whether there is adequate functional cold
	chain capacity to meet needs for routine immunization, campaigns and new
	vaccine introductions
Related	Temperature Alarm Rate
indicators	Temperature in Range
	 Number of Maintenance Visits, Requests and Repairs
	Cold Chain Equipment Uptime
	Cold Chain Capacity Utilization
	Mean Time to Repair Cold Chain Equipment
	Mean Time to Implement Corrective Action
	Mean Time between Refrigerator Failure

Name	On-Time and In-Full Delivery
Description	 Percentage of deliveries delivered on-time and in-full (OTIF), with OTIF defined as: Order fulfilled: Store can fulfil the complete order (i.e., provide all products and quantities requested) On time: Order is delivered when expected (e.g., on a specific date or within a specified time range) Accurate: The correct products are delivered in the correct quantities (i.e., delivered products and quantities match the delivery note)
Purpose	Used to ensure the store has the ability to meet the needs of lower-level stores, as well as the timeliness and reliability of order deliveries. The indicator can be used to monitor incoming shipments and performance of in-country distribution by the national store or outsourced distributor. Including the indicator in a dashboard can facilitate store management improvements: increased reliability, consistency (client receives product needed each resupply period)
	 and efficiency (reduction in emergency orders). Note that OTIF delivery does not consider damage to products during distribution (e.g., broken vials, VVM stage 3 or 4). Other indicators (such as Closed Vial Wastage or Temperature Alarm Rate) should be used to identify such issues. The following questions can be answered by monitoring this indicator: Are deliveries received during the expected time period?
	 If warehousing and/or delivery services are outsourced, have the third-party logistics providers achieved their agreed-upon/contractual service levels? Are orders correctly picked and packed, in terms of products and quantities? Are orders correctly distributed in terms of products and quantities? Have global procurement service agents and freight forwarders delivered products in-full and on-time?
Performanc e	Efficiency Availability
objective Domain	Distribution Stock management
Full indicator name(s)	% of orders delivered on-time and in-full (OTIF)
Dashboard use level	This indicator is recommended in dashboards used by national and store managers at all levels.
Preconditio ns	 This indicator is relevant in supply chains where: Delivery schedule is in place and date dispatched/received is captured Client knows the amount and/or expected amount Stores deliver supplies to lower level stores or facilities (outbound delivery)
System design	 The indicator is relevant for these supply chain systems: Push system with fixed quantities Pull system with delivery

needed		rder req	uested b	y produc	t and qua	antity				
	• C	order pick	ked and c	lispatche	ed by pro	duct and	quantity	,		
	• S	cheduled	l delivery	date or	delivery	range				
	• P	roducts,	quantitie	es and tir	ne of rec	eipt for d	lispatche	d orders	by order	-
Data	• 0)rder deli	very not	е						
sources	• S	ubmitted	l requisit	ion/orde	er					
	• P	roof of d	elivery							
	• D	elivery s	chedule							
	• V	accine a	rival rep	ort						
	• A	dvanced	shipmer	nt notifica	ation					
Data	Data for	this indic	ator is to	be colle	ected and	l compile	d by the	store res	sponsible	for
collection	fulfilling	the orde	rs. If the	data coll	ection sy	stems ar	e manua	l, sampli	ng or sen	tinel site
method	can be u	sed to co	llect data	a for calc	ulation o	f OTIF. If	the sam	ple is lar	ge enoug	h, this
	method	will give a	a good pi	icture of	the actu	al perforr	nance of	the syst	em.	
Calculation	% of ord		ered on-t	ime and	in-full =	# orders	delivere	d OTIF/t	otal # ord	ders
	delivered	d) x 100						-		
	For store						cesses, a	n interm	iediate in	dicator
	(such as	% of on-t	ime deliv	veries) ca	an be use	ed.				
	Example			1						
	Consider a regional store that picks, packs and delivers to four district stores of						nce a			
	month. The date of delivery, scheduled delivery date and data on quantities ordered, dispatched and received were collected from the relevant data sources and compared									
						very date	e and dat	a on qua	antities o	rdered,
	dispatch	ed and re	eceived w	vere colle	ected fro	very date m the rel	e and dat	a on qua	antities o	rdered,
		ed and re	eceived w	vere colle	ected fro	very date m the rel	e and dat	a on qua	antities o	rdered,
	dispatch	ed and re	eceived w	vere colle	ected fro	very date m the rel	e and dat	a on qua	antities o	rdered,
	dispatch	ed and re f orders	eceived w were del	vere colle	ected fro n-time ar	very date m the rel id in-full.	e and dat	a on qua ta sourc	antities o es and co	rdered,
	dispatch	ed and re f orders	eceived w were deli	vere colle	ected fro n-time ar	very date m the rel id in-full.	e and dat evant da	a on qua ta sourc	antities o	rdered,
	dispatch	ed and re f orders ested	eceived w were deli	ntities ived ou	ected fro n-time ar	very date m the rel id in-full.	e and dat evant da leq:-	a on qua ta sourc	es and co	rdered, ompared
	dispatch	ed and re f orders	eceived w were deli	vere colle	ected fro n-time ar	very date m the rel id in-full.	e and dat evant da	a on qua ta sourc	antities o es and co	rdered,
	dispatch	ed and re f orders ested	eceived w were deli	ntities ived ou	ected fro n-time ar	very date m the rel id in-full.	e and dat evant da	a on qua ta sourc	es and co	rdered, ompared
	dispatch identify i	ed and re f orders rednested 30	oreived wwere deli backed 28	vere colle ivered or received 28	ected fro n-time an delivery 1–5	very date m the rel d in-full. of receibt	e and dat evant da fulfilled? No	a on qua ta sourc accruate Yes	antities o es and co time; No	ondered, ompared delivery No
	dispatch identify i Store A	ed and re f orders rednested	Quantities packed packed	Quantities inceceived received	ected fro n-time an Cuped Alivery 1–5 Nov.	very date m the rel d in-full. of receipt 7 Nov.	e and dat evant da fulfilled?	a on qua ta sourc accnrate;	order on- es and co time;	odered, ompared delivery
6	dispatch identify i Store A Store	ed and re f orders understed 30 30	eceived w were deli backed 28 23	vere colle ivered or uantities 28 20	ected fro n-time an Vaaviaa 1–5 Nov. 10–15	very date m the rel d in-full. of receipt 7 Nov. 10	e and dat evant da Unffilled No No	a on qua ta sourc order yes No	ntities o es and co order ou- time; No Yes	ndered, ompared OTIE No No No
Q	dispatch identify i Store A Store B	ed and re f orders rednested 30	oreived wwere deli backed 28	vere colle ivered or received 28	ected fro n-time an Augustation Schednled 1–5 Nov. 10–15 Nov.	very date m the rel d in-full. Vectnal date of Leceipt 7 Nov. 10 Nov.	e and dat evant da fulfilled? No	a on qua ta sourc accruate Yes	antities o es and co time; No	ondered, ompared delivery No
	dispatch identify i Store A Store B Store	ed and re f orders of Grantities 30 30 23	eceived w were deli backed 28 23 23 23	vere colle ivered or Gnantities 28 20 23	ected fro n-time an Logo Chedraled Logo Chedraled L	very date m the rel d in-full. Vctnal date of receipt 7 Nov. 10 Nov. 12	e and dat evant da Urder No No Yes	a on qua ta sourc Guder Yes No Yes	ntities o es and co order ou- times No Yes Yes	ndered, ompared JU O U O U O I V o No Yes
	dispatch identify i Store A Store B Store C	ed and re f orders understed 30 30	eceived w were deli backed 28 23	vere colle ivered or uantities 28 20	ected fro n-time an August of the sector nov. 10–15 Nov. 10–15 Nov.	very date m the rel d in-full. Vetnal date of Leceipt 7 Nov. 10 Nov. 12 Nov.	e and dat evant da Unffilled No No	a on qua ta sourc order yes No	ntities o es and co order ou- time; No Yes	ndered, ompared OTIE No No No
	dispatch identify i Store A Store B Store C Store	ed and ref f orders of antities 30 30 23 15	eceived w were deli and the second second and the second second and the second second and the second	vere colle ivered or 0nantities 28 20 23 15	ected fro n-time an Apply of the second second from the second second second from the second second second from the second second second second from the second second second second second from the second second second second second second second second second second second second second second second second	very date m the rel d in-full. Vetral date 7 Nov. 10 Nov. 12 Nov. 13 Nov.	e and dat evant da Voder No No Yes Yes	a on qua ta sourc 	ntities o es and co - - - - - - - - - - - - - - - - - - -	rdered, ompared JU Ho No No Yes Yes

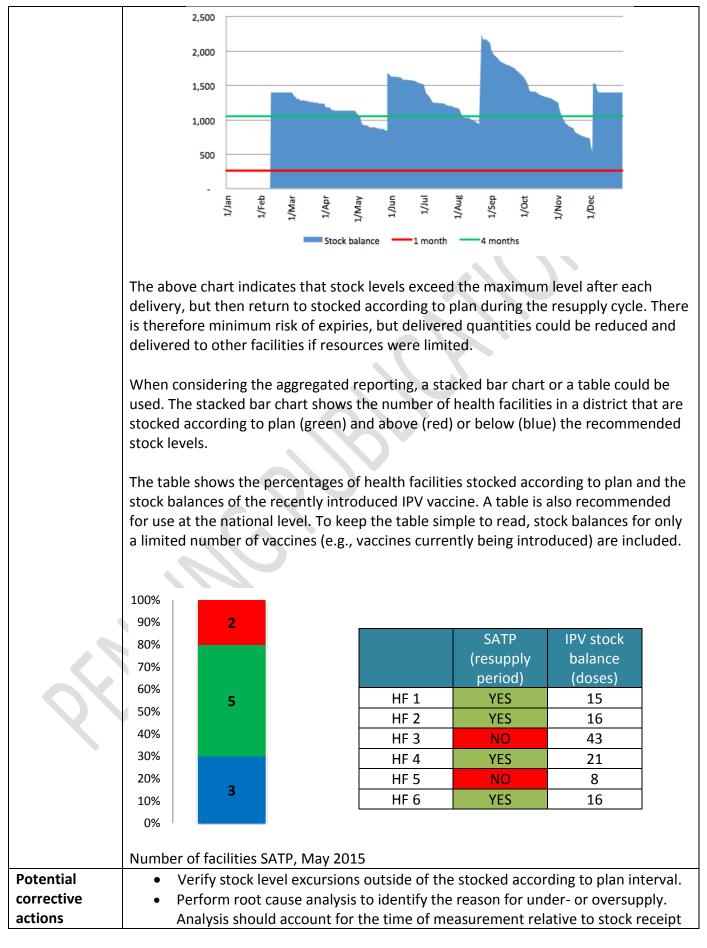
	The National Logis deliveries and nee quarter. The following tabl delivered on-time	ds to aggregate r e shows the num and in-full during	ber of each is the months	nation regiona s of the	al store perform al store's total d e first quarter.	mances for the deliveries that	-
	Store	January	Februar	y	March	Q1	
	Regional store	2 of 4	2 of 5		3 of 6	47%	
	1 Regional store	(50%) 4 of 6	(40%) 5 of 8		(50%) 6 of 8		
	2	(67%)	(63)		(75%)	68%	
	Regional store	3 of 3	3 of 4		4 of 4		
	3	(100%)	(75%)		(100%)	92%	
		1 of 3	2 of 3		2 of 3		
	National store	(33%)	(67%)		(67%)	56%	
	Average store	10 of 16	12 of 20)	15 of 21		
	OTIF	(63%)	(60%)		(71%)	65%	
Visualizatio n and interpretati	On-time and in-ful less than 50% to n The National Logis regional stores for visualize and mana	nore than 91%. tics Working Gro the past quarter	up (NLWG) i . The NLWG	s revie had es	wing the perfo	rmance of the	three
on						0715	
	OTIE < COV (oritic	rat rad)			Stores	OTIF	
	OTIF <60% (critic OTIF >60% and <		llow	Nat	ional store	average	
	OTIF >80% (norn		110 W)		onal store A	95% 78%	
		iai, green)	F		onal store B	88%	
					onal store C	58%	
	100%						
	90% 80%		× •	R	egional Store A		
	70%				egional Store B		
	60%				egional Store C		
	F.0%/				•		
	50%				ational Store		
	40%						
	30% January	February N	March				
	Regional store C h disaggregated OTI			•		•	

	Q1	March
	100% 90% 80% 70% 60% 50% 40% 30% 20% 10% 0%	 Dispatch & Distribution Inaccurate Order Not Fulfilled On-Time Arrival Missed On-Time Arrival Missed
Potential corrective actions	 Improve or define Revise demand pla If services are outs distribution service Adjust delivery sch transportation serv Improve forecastin supplying stores Negotiate with pro shipments to the c Assess system or p 	g and procurement procedures to ensure adequate stock at curement service agents and freight forwarders on in-bound ountry olicy changes (e.g., outsourcing or changing distribution system) se inventory policies including buffer stock and minimum and
Related indicators	 On-Time Arrival In-Full Arrival In-Full Dispatches On-Time Dispatches Order Accuracy % of Deliveries with 	25

Name	Stocked According To Plan
Description	This indicator measures the percentage of health facilities or stores maintaining appropriate (as defined by local policies) levels of vaccine and immunization product stock during a particular time frame, as compared to the overall number of facilities in the area. Stocked according to plan (SATP) is defined as stock levels between set minimum and maximum levels.
Purpose	Used to monitor and manage immunization products and as a warning to avoid stock- outs or wastage. Diversions from the planned stock levels can signal risk of stock-outs (if significantly below the minimum level) or closed vial wastage (if significantly above the maximum level). For stores, the indicator performance provides information on the ability of the store to dispatch the products and quantities needed by the health facilities.
	 The following questions can be answered by monitoring this indicator: Is there a risk of stock-outs? Is there a risk of overstock and expiry? Will the supplied quantities be enough until next delivery? Are the demand methodology and assumptions adequate? Are the inventory policies and practices adequate?
Performance	Availability
objective	Efficiency
Domain	Stock management
Full indicator	 % of health facilities stocked according to plan
name(s)	 % of districts with x% of facilities stocked according to plan
	 % of stores stocked according to plan
Dashboard use	This indicator is recommended in dashboards used by national and sub-national
level	managers.
Preconditions	 This indicator is relevant in supply chains where: There are established minimum and maximum levels for products for each health facility and store. Minimum stock level is considered the safety stock that is different from the reorder stock level. The maximum stock level is the safety stock plus the expected consumption between deliveries.
System design	Relevant for supply chain systems with minimum stock level equal to safety stock. The indictor is not relevant in systems where the minimum stock level is considered equal to the reorder level, as the stock is expected to go below the minimum stock level. In these systems, alternative indicators such as Full Stock Availability and Closed Vial Wastage may be better employed.
Data needed	Stock balance
	 Minimum and maximum levels
Data sources	 Stock cards/ledgers Physical inventory count Logistics management information system (LMIS)
Data	Stock balances should be collected at least twice per resupply period: just after and
collection	before delivery, to provide the highest and lowest stock balances in the resupply
method	period.

Calculation	hand) to the estab have stock balance according to plan maximum stock le regarding the num	to plan is determined l blished minimum and m es below, within or abo occurs when the stock l vels, which are typically bber of months of stock	aximum levels to ident ve the recommended l balance is between the y set by national policy to be held in each type	tify which products levels. Stocked e set minimum and , for instance e of store or facility.			
	In a store or health facility, each product can be assessed as stocked accordin plan. Alternatively a set of tracer products can be considered. When aggrega indicator at higher levels, then a health facility or store is considered stocked according to plan if all vaccines and immunization supplies are stocked accord plan.						
	•	cked according to plan (to plan for all or a set o					
		es stocked according to set of tracer products),					
	MinimumMaximum	, the inventory policy fo level: 50 doses level: 100 doses ne facility's actual stock od.		ning and near the end			
	Vaccine	Stock balance (start of supply period, doses)	Stock balance (end of supply period, doses)	Stocked according to plan for this antigen			
	Rota	160	44	NO			
	PCV	93	63	YES			
	Penta	87	56	YES			
	OPV	75	53	YES			
	Measles	109	48	NO			
	IPV	83	43	NO			
	The health facility	cked according to plan = would be considered n cked within minimum a	ot stocked according to	•			
	of each month. Th	n health facilities that r e district has quarterly er products are used to	resupplies, so the indi	cator is reported			

			July	August	September	SATP in Q3	
		HF 1	SATP	SATP	BELOW	NO	
		HF 2	SATP	SATP	SATP	YES	
		HF 3	ABOVE	ABOVE	ABOVE	NO	
		HF 4	ABOVE	SATP	SATP	NO	
		HF 5	SATP	SATP	SATP	YES	
		HF 6	SATP	SATP	BELOW	NO	
		HF 7	SATP	SATP	SATP	NO	
	SA	ATP (monthly)	71%	86%	57%	29%	
Visualization and interpretation	% health The mon that were after sup of the tra resupply The stock easier int levels are track the predicts	facilities SATP (facilities SATP (thly calculation e stocked accor plies were rece acer products by period, only 29 k balances in sto terpretation of t e included in the stock levels, wh the future stock	Q3) = 2/7 x 10 s show that by ding to plan in ived) reach be y the end of th <u>% of health fa</u> ores can be vis the SATP visua e graphs. In he hereas in the r	0 = 29% the end of the the beginning low the minim re resupply per cilities were st ualized in char lization, the m ralth facilities, national store, on consumptio	of the resupp ium stock leve riod. Overall fo ocked accordi rts, such as the ninimum and n a simple graph a sophisticate	oly period (righ of the quarterly ng to plan. two below. Fo naximum stock n can be used t d graph that	t nore y or <
	0		Defailed overview of	BCS vacuum availability, as of April	17, 2018		
		1.600,000	1				
		1,400,000				~	
		1.980.000					
		1,00,000 800,000 800,000	- and		·		
0		2000					
	100				- Concernance (Concernance)		
		for Vaccines (Vi					



	 (i.e., stock levels should be at or slightly exceed the maximum upon stock receipt and decrease over time). Prioritize actions for critical or problematic products and/or locations with low stocked according to plan percentages. Review and revise inventory and distribution policies including minimum and maximum levels.
Related	Full Availability
indicators	Functional Status of Cold Chain Equipment
	On-Time and In-Full Delivery
	Closed Vial Wastage
	Cold Chain Capacity Utilization

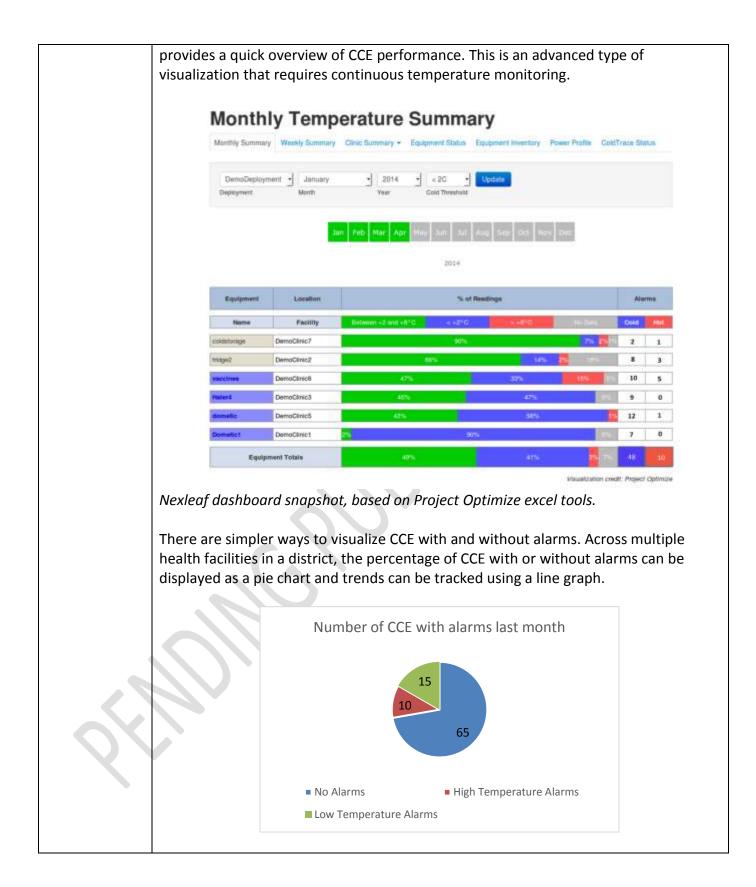
Name	Temperature Alarm Rate
Description	Number of times the temperature inside cold chain equipment (CCE) exceeds or drops below a reference range. The indicator is applicable where vaccines are stored and during transportation. CCE is defined as all refrigerators, freezers, passive storage devices, and walk-in cold rooms and freezer rooms designated for storing vaccines.
Purpose	Used as a proxy for measuring vaccine potency and safety. Exposure to temperatures outside this range indicates a risk of heat or freezing damage to sensitive vaccines.
	 The following questions can be answered by monitoring this indicator: Is there risk of heat exposure to vaccines? Is there risk of freeze damage to vaccines? Is cold chain equipment functioning properly? Which CCE devices are in need of repair or replacement?
Performance objective	Potency
Domain	Cold chain management
Full indicator name(s)	 Rate of heat and cold alarms per monitoring period (e.g., per month) Number of CCE devices with more than a certain number of temperature alarms during a monitoring period
Dashboard use level	This indicator is recommended in dashboards used by sub-national and national managers. Visual monitoring of temperature (i.e., through monitoring of 30-day temperature recorders and/or thermometers) is recommended in health facilities and stores.
Preconditions	The indicator is relevant for all types of immunization supply systems, for all locations where immunization products are stored. A mechanism for routinely measuring and recording temperature is needed in each device designated for storing vaccines.
System design	Relevant in all types of logistics systems.
Data needed	Continuous or point-in-time temperature readings recorded over a time period. Continuous temperature monitoring is highly preferred, since it allows greater accuracy in detecting temperature fluctuations. For primary and sub-national stores, programmable electronic temperature and event logger systems are the best option. In smaller stores and health facilities, 30-day electronic temperature records with a stem thermometer as a backup are considered best practice. A stem thermometer alone only indicates the temperature at the time a reading is taken, which is no more than 14 times per week. A 30-day temperature logger takes at least a thousand readings per week. ²
	The number of excursions, or alarms outside the designated temperature ranges, is needed. Alarm thresholds are set by WHO:

² World Health Organization, *How to Monitor Temperatures in the Vaccine Supply Chain: WHO vaccine management handbook – Module VMH-E2-01.1, WHO, Geneva, 2015, <http://apps.who.int/iris/bitstream/10665/183583/1/WHO_IVB_15.04_eng.pdf?ua=1>, accessed 7 November 2015.*

[
	 An excursion is defined as any event during which the temperature inside the cold chain equipment goes below 2° C or above 8° C. A high temperature alarm is defined as any event during which the temperature goes above 8° C for 10 continuous hours. A low temperature alarm is defined as any event during which the temperature goes below -0.5° C for one hour. For locations measuring and recording temperatures manually twice daily, alarms may be difficult or impossible to record. Record any excursion outside the range of 2°C to 8°C for refrigerators and -15°C to -25°C for freezers. A point-in-time 'temperature in range' indicator may be used instead. However, a point-in-time temperature reading within temperature range does not provide any indication about temperature excursions that may have occurred at other times throughout the day when the
	temperature was not being recorded (e.g., a cold exposure overnight, when ambient
	temperatures dropped). Note that WHO no longer recommends stem thermometers
	and point-in-time recording of temperature as the primary means to monitor
	temperature in cold chain equipment. ³
Data sources	 Continuous temperature recording devices, including 30-day temperature recorders. Wherever possible, temperatures should be recorded automatically. High/low temperature alarms (built into CCE or temperature monitoring devices)
	 Proof of delivery (POD) for measuring temperature during transit if a temperature groupling during is included.
Coloulation	temperature recording device is included
Calculation	Temperature alarm rate = number of high and low temperature alarms per reporting period This indicator can also be calculated using the number of CCE devices with more than
	a set threshold of temperature alarms in a given period.
5	It can be further broken down by reasons for alarms (if known) or into 'resolved' and 'unresolved' alarms. That is, an alarm due to a resolved power outage would not be treated the same as an alarm due to mechanical problems. The alarm rate can also be disaggregated by facility, by device or by device type (make, model, energy source, etc.) to monitor performance.
	<i>Examples</i> A facility has one ice-lined refrigerator with a 30-day temperature logger. During a supervisory visit, the temperature data is downloaded, and the following temperature alarms are noted:

³ World Health Organization, 'The Vaccine Cold Chain', Module 2 in *Immunization in Practice*, WHO, Geneva, p. 22, <www.who.int/entity/immunization/documents/iip2014mod2aug4.docx?ua=1>, accessed 7 November 2015.

	Data: April 2 Data: April 20]	
	Date: April 3 Timo: 04:15			Date: April 29					
	Time: 04:15				Time: 16:34				
	Temp: -1.2				Temp: 12.3				
	Alarm: COLD			Alarm: HEAT					
	Durat	ion: 1h 24n	nin	Duration: 14h 06min					
	This facility had an alarm rate of 2 alarms per month during the month of April. Alternatively, heat and cold alarms can be reported separately, with an alarm rate of 1 alarm per month for each. If the cause of alarms is known, this indicator can be further disaggregated (e.g., 1 heat alarm due to power outage). In a district comprising 40 health facilities, each of which is using 30-day temperature							rate of 1 De	
	recorders, there were a total of 16 alarms during the past month; 4 high temperature alarms and 12 low temperature alarms. The rate is reported as 4 high temperature alarms per month and 12 low temperature alarms per month.								
Visualization	Continuous temperature recording devices can provide tabular readouts of								
and	temperature data, including alarms.								
interpretation									
•		Lower alarm limit							
		Date	Average temperature	Status	Minimum temperatu re	Duration out of range	Alarm trigger time		
		02.12.2014	+ 4.5 C	ok	+ 4.1 C				
		02.12.2014	+ 4.3 C	ok	+ 4.0 C			-	
		30.11.2014	+2.2 C	ALARM!	-1.5 C	2h 30min	05:05		
		29.11.2014	+3.4 C	ok	+2.5 C	211 3011111	03.03	_	
	Across facilities, the alarm rate can be displayed in a colour-coded table to highlight facilities with frequent temperature excursions. These may be targeted for repair or replacement of CCE or for additional training on vaccine management.								
				Alarm	Alarm	Alarm			
			Facility	rate –	rate –	rate –			
				June	July	August	t		
			Health facility A	0	1	0			
			Health facility B	0	0	0			
			Health facility C	4	3	5			
			Health facility D	1	0	0			
	Listing or visualizing only poorly performing fridges or facilities can allow for easier prioritization of facilities that need immediate attention. The overview below shows the alarms for six CCE devices over a month (January). The								
	graph highlights the high and low temperature alarms and the percentage of time								
	that the CCE was within the recommended temperature range during the month. It								



	$\begin{array}{c} 90\\ 80\\ 70\\ 60\\ 50\\ 40\\ 90\\ 20\\ 10\\ 0\\ 0\\ 10\\ 0\\ 0\\ 10\\ 10\\ 0\\ 0\\ 10\\ 1$						
	considerations related to the value of vaccines at risk should be taken into account.						
	For instance, more urgent action might be needed for a walk-in cold room storing thousands of doses of vaccines than for a single refrigerator at health facility level.						
Potential corrective actions	 Ensure that facilities follow standard operating procedures through supportive supervision. For instance, facility staff should remove vaccines from CCE not maintaining temperature within recommended ranges in accordance with contingency plans and should discard vaccines that have VVM stage 3 or 4 and vaccines that fail the shake test. Determine cause of equipment dysfunction; solicit repair or replacement of non-functional equipment Ensure that contingency plans are in place for all facilities Perform regular routine maintenance of all CCE to prevent future breakdown Train facility staff to improve inventory management practices Use temperature alarm profiles of various types and models of CCE to inform procurement Use temperature alarm profiles to plan for repair and replacement of CCE Cold Chain Equipment Uptime Cold Chain Equipment Uptime Cold Chain Capacity Utilization Mean Time to Repair Cold Chain Equipment 						
Related indicators							
	 CC Energy Source Report Number or % of Vaccines Discarded Due to Heat Exposure or Freeze Damage 						