



# **UNIVERSAL IMMUNIZATION** **PROGRAMME WITH** **RECENT ADVANCES**



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# Universal Immunization Programme

## (Scope and Scale)

**Annual target**

**2.67 crore newborns;  
2.9 crore pregnant women**

**Vaccine against VPDs**

**9 nation wide;  
3 sub-nationally (JE, Rota, PCV)**

**One of the largest Public Health Programmes**

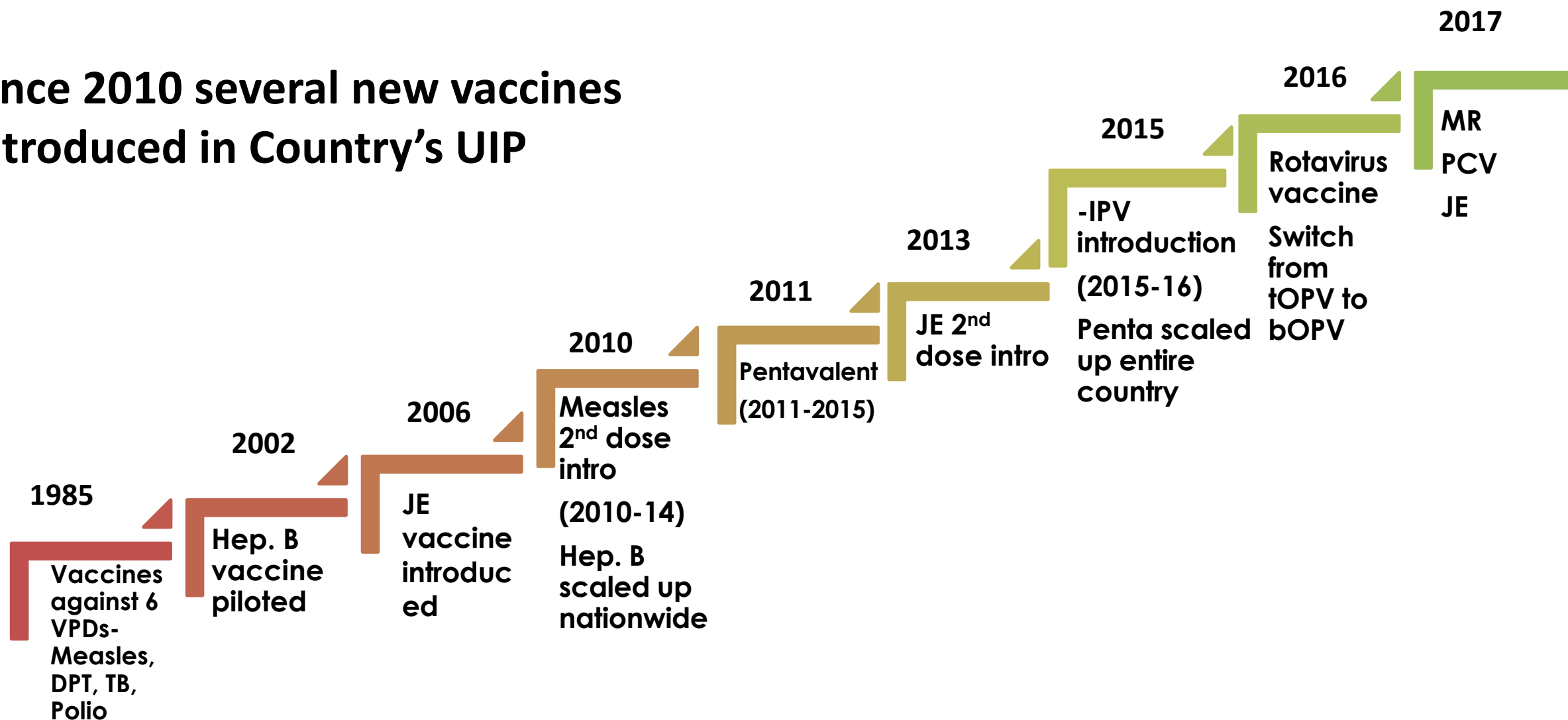
**≈1.2 crore sessions planned per year**

**~29,000 cold chain points for storage  
and distribution of vaccines**

***Make in India:* Largest vaccine manufacturing capacity in the world**

# Roadmap of vaccine Introduction

Since 2010 several new vaccines introduced in Country's UIP



# Rapidly changing landscape of Universal Immunization Programme

## Milestones

- 2014: India declared Polio free
- 2015: Maternal & Neonatal Tetanus Elimination validation

## Improving Coverage

- 2015: Mission Indradhanush
- 2017: Intensified Mission Indradhanush
- 2018: Gram Swaraj Abhiyan (GSA)/Extended GSA

## Improving Quality

- NCCRC / NCCTC
- EVM assessment
- eVIN expansion
- Capacity building of HR

## New vaccines introduced

- 2015: Inactivated Polio Vaccine (IPV)
- 2016: Rotavirus Vaccine (RVV)
- 2017: Measles-Rubella (MR) Vaccine and Pneumococcal Conjugate Vaccine (PCV)

# Two milestones achieved



On 27<sup>th</sup> March 2014, South-East Asia Region of WHO, including India, certified POLIO-FREE



On 14<sup>th</sup> July 2016, WHO certified India for eliminating maternal and neonatal tetanus



## Why an Immunization program?

While a health worker giving a vaccine in a clinic to one child may seem a simple procedure, a well organized public health program is needed to ensure vaccines are given to several thousands of children. The Immunization program is an important public health program of most government health departments and other health agencies.

### Features of the program:

- It is based on scientific evidence and rationale.
- It needs an effective management system to ensure vaccine quality and consistent delivery.
- It requires well trained and motivated personnel to provide regular dependable services.
- It requires reach and continued acceptance among beneficiaries
- With adequate coverage and quality it results in prevention of the diseases in the community.

Effective management systems for the immunization program need to consider aspects of

- resource management including human, infrastructure and financial resources
- supply chain management,
- personnel management including capacity building,
- social mobilization including effective community participation and
- data review mechanisms such as authentic reporting and surveillance.

Health programs such as the Immunization program also needs to be linked to other health and development programs that use the same health systems, infrastructure and personnel.

## What are various immunization related programs in India?

**UIP:** The Universal Immunization Programme (UIP), launched in 1985 to progressively cover the country, aimed to reduce mortality and morbidity from the six vaccine preventable diseases (measles vaccine was added in 1985). Under the program, indigenous vaccine production capacity was enhanced and a national cold chain established.

Vitamin A supplementation was included in 1990. Within the Universal Immunization program, newer vaccines such as Hepatitis B are being introduced country-wide from 2011. Japanese Encephalitis vaccine is a part of the routine immunization package in endemic districts while Penta-valent vaccine is being introduced in the states of Kerala and Tamil Nadu.

**NRHM:** Launched in 2005, the National Rural Health Mission has made way for a comprehensive and effective integration of health programs including Immunization. It also provides for resources, mechanisms and flexibility for effective program management. Additional manpower for management and linkages with the villages called ASHA (Accredited Social Health Activist) has helped boost health program implementation and acceptance. As a part of the integrated approach, immunization sessions are being used as a platform to provide other health, nutrition and sanitation related services and these special days are called Village Health and Nutrition Days (VHNDs).

**Polio NIDs and SNIDs:** With Polio eradication as a goal, states in India have few to several rounds of Polio supplementary Immunization activities. These are called National (NID) and sub-national Immunization days (SNID) and consist of booth based and house-to-house vaccination activities lasting several days.

**Measles catch-up and second opportunity:** Measles Mortality reduction is the goal of this activity where a second opportunity for measles vaccination is being given to children either through a catch-up activity or through introduction in the regular immunization schedule.

## **What are vaccine preventable diseases?**

Vaccines are now available to prevent certain diseases. In most cases if a potent vaccine is given correctly, that is at appropriate time, dosage and technique, it can prevent the disease from occurring in a vaccinated individual despite his exposure to the disease. Diseases for which an effective vaccine has been made and is available for use are known as a Vaccine Preventable Disease (VPDs).

## **What diseases are prevented through vaccines used in the Universal Immunization Program (UIP)?**

Presently, the Universal Immunization Program in India provides vaccines mainly to children below 5 years of age and pregnant women for the following vaccine preventable diseases:

1. Tuberculosis
2. Poliomyelitis
3. Diphtheria
4. Pertussis (whooping cough)
5. Measles
6. Tetanus
7. Hepatitis B
8. Japanese encephalitis (in endemic districts)

Vitamin A is not a vaccine, but a nutritional supplement which prevents many deficiency related conditions. However administration of Vitamin A is also a part of the Universal Immunization Program.

Some other diseases have combined vaccines so as to avoid multiple shots, for example DPT for Diphtheria, Pertussis and Tetanus. This is also called a triple antigen. A penta-valent vaccine (5 vaccines together) is also being considered for introduction in the UIP. This will include DPT+ Hepatitis B vaccine+ vaccine for Haemophilus B.



## 1. Tetanus:

**Vaccine: Tetanus toxoid:** packaged as liquid in 10 dose vial. (The vaccines DT, DPT, DTaP also protect against tetanus but here tetanus toxoid is in combination with other antigens)

### Disease:



#### Tetanus

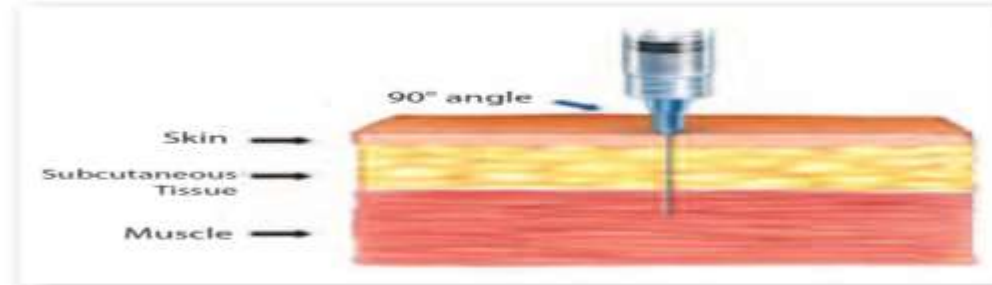
Suck & cry in first 2 days of life, illness between 3-28 days of life, inability to suck followed by stiffness of neck & body &/or jerking of muscles

### Dose:



Dose: 0.5 ml; Vial: 10 doses

### Route:



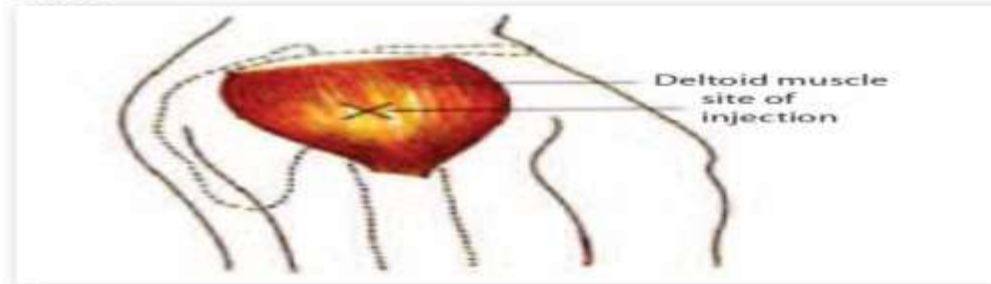
Intramuscular Injection

### Schedule:



**Early in pregnancy and at least 4 weeks later**  
For subsequent pregnancy within 3 years of last where 2 doses of TT were given, here only 1 booster shot required.

### Site:



Deltoid Muscle

### Method:



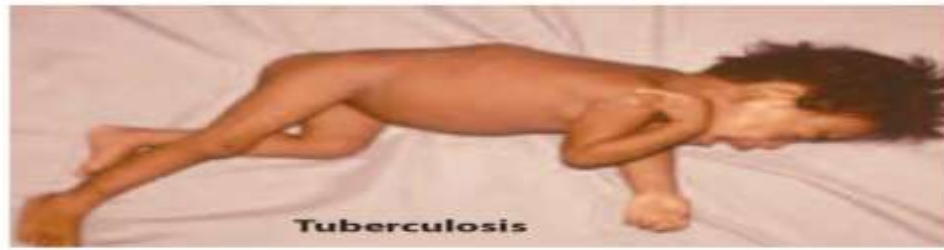
Stretch skin flat between finger and thumb on either side of injection site

Remember: T series vaccines are destroyed if frozen and should be stored and transported in temperatures between 2 and 8 degrees Celsius.

## 2. Tuberculosis:

**Vaccine:** BCG (Bacillus Calmette-Guérin) powder in a 10 dose amber vial reconstituted with Normal saline

**Disease:**



Tuberculosis

\*history of contact with suspected/confirmed case of pulmonary TB  
\*Weight loss, cough and wheeze not responding to ARI antibiotics

**Dose:**



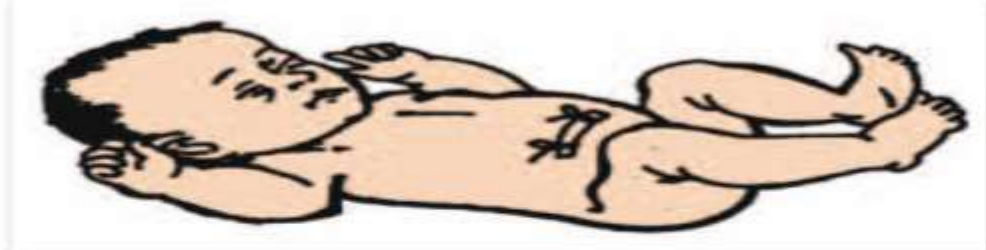
Dose: 0.1 ml (0.05ml for less than 1 month);  
Diluent: 1 ml sodium chloride; Vial: 10 doses

**Route:**



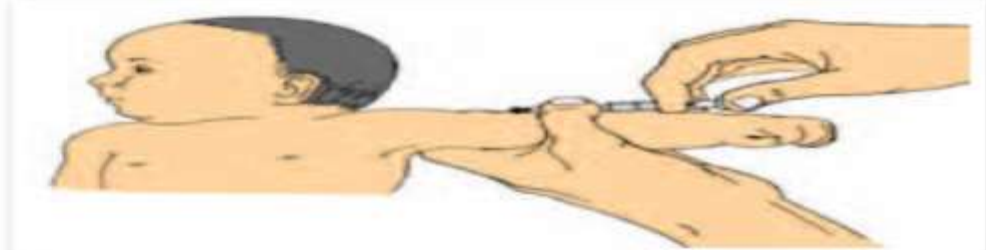
Intra-dermal Injection

**Schedule:**



Give at birth or as early as possible in the first 12 months.

**Site:**



Left Upper Arm

**Method:**



Gently pull skin under arm to stretch skin at injection site

Remember: after BCG vaccination, the skin gets raised and later a scar will form.

However, BCG vaccination is not to be repeated even if the scar does not form. BCG if missed earlier can be given till 1yr of age of the child. After reconstitution, BCG should be used within 4 hours.



### 3. Poliomyelitis:

**Vaccine:** Oral Polio vaccine (Trivalent: Sabin) liquid in a 20 dose vial with dropper

**Disease:**



\*Sudden weakness & paralysis of leg(s), &/or arm(s) &/or trunk

\*Paralysis not present at birth or due to serious injury/mental retardation

**Dose:**



**Dose: 2 drops; Vial: 20 doses**

**Route:**



**Put two drops directly in mouth of child**

**Schedule:**



**Zero Polio dose at birth till 15 days, 11/2 months....(14 weeks) and at 16 to 24 months with DPT booster"**

**Site:**



**Mouth**

**Method:**



**Oral administration**

Remember: During Polio supplementary rounds, other types of Polio vaccines may be used. These may be mono-valent OPV, bivalent OPV or even Injectable Polio Vaccine (IPV) These should be distinguished from the trivalent OPV to be used during UIP sessions.

#### 4. Diphtheria, Pertussis and Tetanus:

**Vaccine:** DPT (Trivalent) liquid in a 10 dose vial for injection

**Disease:**



**Diphtheria:** Sore throat with gray patch(es) in throat  
**Pertussis (whooping cough):** repeated & violent coughing, with: cough persisting for 2+ weeks, fits of coughing, cough followed by vomiting, typical whoop in older infants.

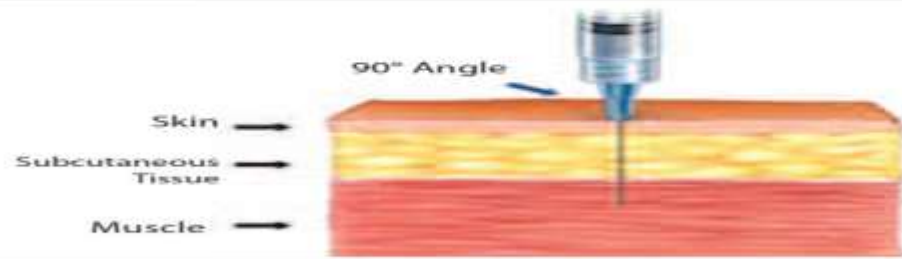
**Tetanus:** suck & cry in first 2 days of life, illness between 3-28 days of life, inability to suck followed by stiffness of neck & body &/or muscle jerking

**Dose:**



**Dose: 0.5 ml; Vial: 10 doses**

**Route:**



**Intramuscular Injection**

**Schedule:**



**Birth 24 hours of following delivery, 1½ mths (6 wks), 2½ mths (10 wks), 3½ mths (14 wks)**

**Booster at 16-24 months and at 5 years**

**Site:**



**Outer Mid-thigh (Antero-lateral side of mid-thigh)**

**Method:**



**Stretch skin flat between finger & thumb on both sides of injection site**

Remember: Giving DPT in buttocks may injure the sciatic nerve and cause paralysis. It should never be given there. Instead, it should be given in the outer mid-thigh. Between two doses of DPT there should be a gap of at least 4 weeks. DPT boosters are to be given at 16-24 months and in the 5th year.



## 5. Hepatitis B vaccine

**Vaccine:** Hepatitis B vaccines: packaged as liquid in 10 dose vial.

**Disease:**



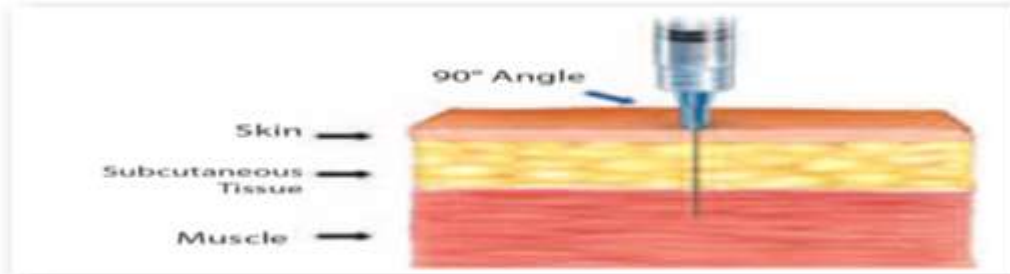
Infection of the liver causing yellow discolouration of skin and mucous membrane, sometimes lead to severe complications like liver failure or chronic disease carrier.

**Dose:**



**Dose:** 0.5 ml; **Vial:** 10 doses

**Route:**



**Intramuscular Injection**

**Schedule:**



at birth within 24 hours of delivery, 1½ mths (6 wks), 2½ mths (10 wks), 3½ mths (14 wks)

**Site:**



**Outer mid-thigh (Antero-lateral side of mid-thigh)**

**Method:**



**Stretch skin flat between finger & thumb on both sides of injection site**

**Remember:** Hepatitis B vaccines are very cold sensitive and are destroyed if frozen and should be stored and transported in temperatures between 2 and 8 degrees Celsius.



## 6. Measles:

**Vaccine:** Measles containing vaccine: packaged as powder in 5 dose amber colored vial. Needs reconstitution with a diluent which is: pyrogen free double distilled water.

### Disease:



**Measles:** fever with rash with cough or running nose or red eyes.

### Dose:



**Dose:** 0.5 ml; **Diluent:** 2.5 ml double distilled water;  
**Vial:** 5 doses



**Subcutaneous Injection**

### Schedule:



**9 months completed - 12 months (39 – 52 weeks).**

If a child does not receive Measles before the 12th month, give a dose as soon as possible before 5 years of age.

### Site:



**Right Upper Arm**



**Pinch skin through left index finger & thumb**

**Remember:** Following reconstitution the measles vaccine has to be used within 4 hrs.

A second dose of measles is to be given with DPT booster at 16-24 months of age initially in select states of India and soon (following Supplementary Immunization campaigns) throughout the country.

## 7. Vitamin A:

**Vitamin A** is not a vaccine but an important micronutrient for maintaining normal growth, regulating cellular proliferation and differentiation, controlling development, and maintaining visual and reproductive functions. However, it is included in the Universal Immunization schedule.

**Vitamin A deficiency** (VAD) increases the risk of disease and death from severe infections such as measles and diarrhoea. In young children VAD can also cause growth retardation. VAD affects many tissues in the body; however its effect is most apparent on the eye. Children with clinical VAD face difficulty in seeing in the night termed as 'night blindness'. At a more severe stage, it results in Bitot's spots, Corneal Xerosis/ ulceration, Keratomalacia and Corneal scar.



Keratomalacia, Bitot's spots, Night Blindness. Other sub-clinical symptoms leading to increased severity of disease and mortality



100,000IU/1 ml (9 months completed to 12 mths)  
200,000IU/2 ml (Over 1 year). Give correct doses

**Schedule:** The first dose of Vitamin A is administered at 9-12 months along with measles. The second dose is scheduled with DPT booster at 16 months. Thereafter, 1 dose is to be given every six months till the age of 5 years, that is, a total of 9 doses till the age of 5 years. In many states Vitamin A is also given through Bi-annual campaigns, in which case, the same dose need not be repeated in the routine immunization sessions.



At 9 months with measles, 16 months with DPT booster, then one dose every 5 months up to the age of 5 years



### Oral Administration

(use only spoon provided observing marks for 1 ml and 2ml)

**Route of administration:** Orally, always using the spoon supplied with the Vitamin A bottle.



## 8. Japanese Encephalitis:

**Vaccine:** live attenuated SA 14-14-2 JE vaccine.

Multi-dose vials with 5 doses, supplied with the diluent vial of 2.5 ml which contains Phosphate Buffer solution.

The vaccine should be reconstituted with the supplied diluent only. After reconstitution it turns into a transparent orange red or light pink liquid. After reconstituting the time of reconstitution should be noted on the vial. The reconstituted vaccine *should be used within two hours of reconstitution*, beyond which the vaccine should be discarded.



**Disease:** a person of any age, at any time of the year with sudden onset of fever and a change in mental status (drowsy, confusion, inability to talk, disoriented or coma) and /or convulsions. This usually follows infection by the Japanese Encephalitis virus introduced through a mosquito bite.

**Schedule** In UIP schedule, following campaigns in an endemic district, JE vaccine should be introduced to new cohorts (children who were underage/not born during the campaign) at 16 to 24 months along with DPT Booster.

**Dose:** 0.5 ml

**Site:** left upper arm

**Route:** subcutaneous

**Method:** pinch skin of left upper arm between thumb and index finger and give at 45 degrees

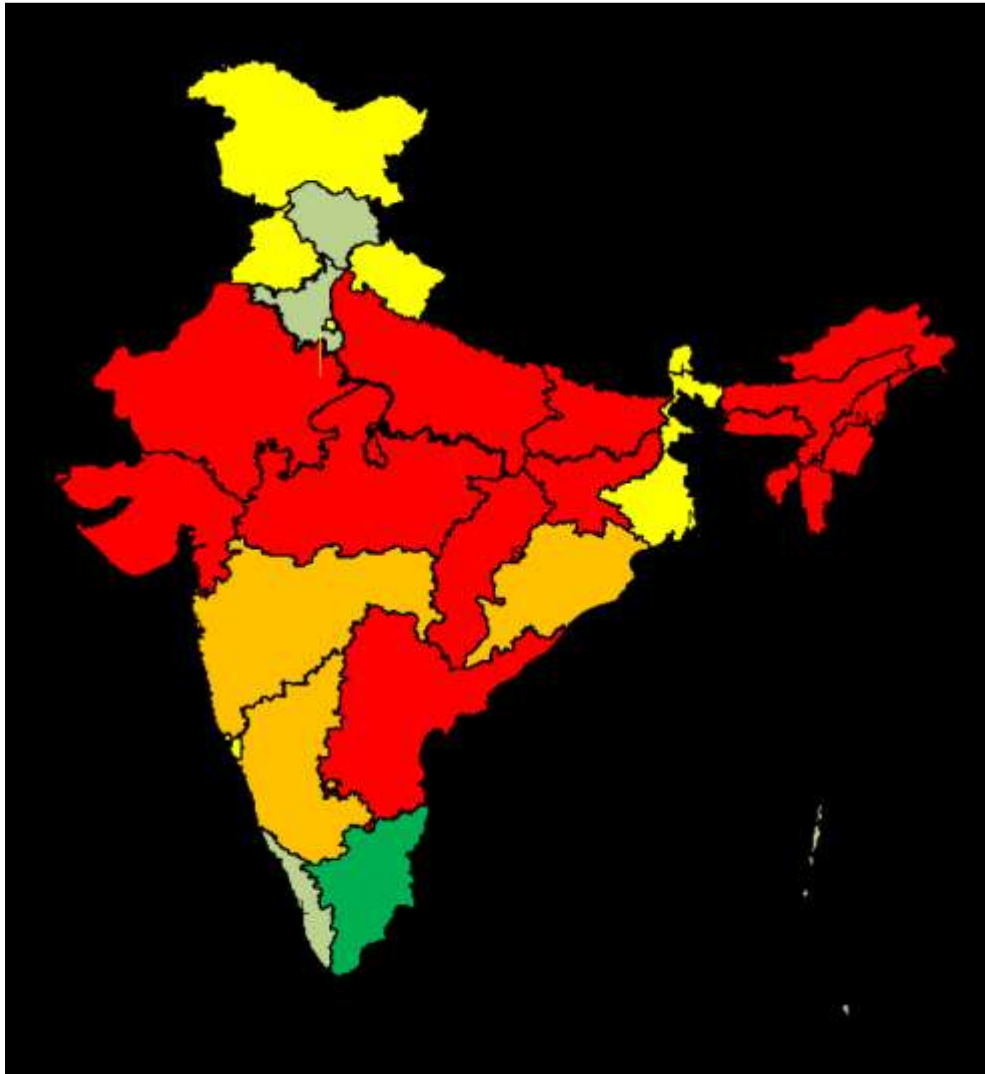
**Contraindications (situations when not to give vaccine):** High Fever (Vaccination to be done only after advise from a Medical officer), Severe malnourishment, Acute infectious disease , Ear infection, Tuberculosis, Heart, liver and kidney problems, Pregnancy , Allergy , Convulsions, Person treated with any immunosuppressive therapy , Person with a proven or suspected hypersensitivity to Kanamycin or Gentamicin.



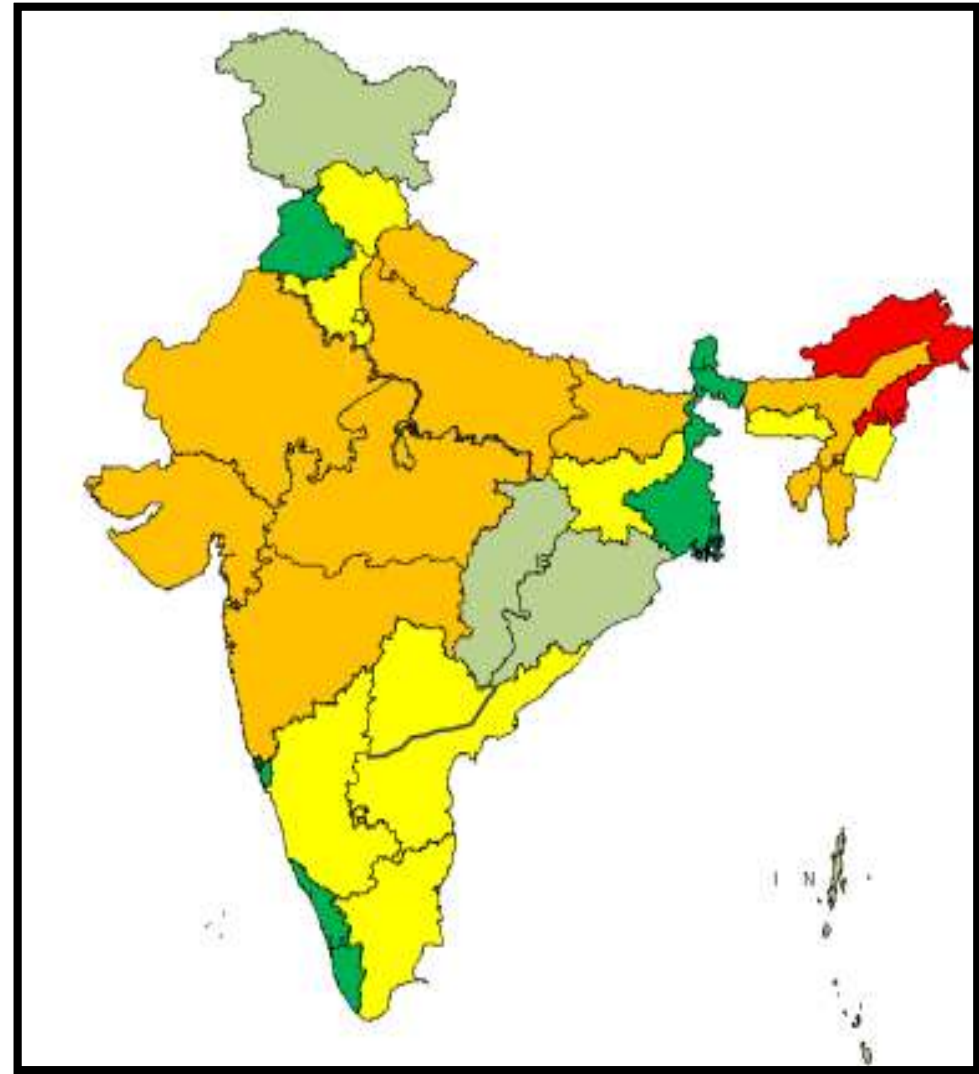
Japanese Encephalitis vaccination is administered under Universal Immunization Program only in selected endemic districts following large scale vaccination campaigns

# Immunization coverage Trends

# Immunization Coverage (FIC)



India: 43.5% ranging from 21% to 81%, NFHS-3, 2005-06



India: 62% ranging from 36% to 91%, NFHS-4, 2015-16



# Mission Indradhanush (MI)



Launched on 25<sup>th</sup> December 2014



- Increasing full immunization coverage to 90% and sustain it through RI
- 554 districts covered in six phases – including Intensified MI
- One of the flagship schemes under Gram Swaraj Abhiyan (GSA) & Extended GSA

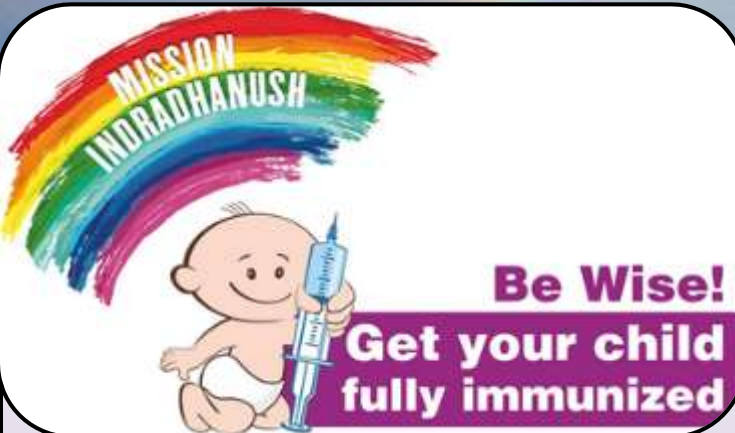
Mission  
Indradhanush  
included under  
PRAGATI

Reviewed by  
Hon'ble Prime  
Minister of India

Impact of MI in  
improving  
immunization  
coverage  
acknowledged

However,  
national coverage  
target of 90% not  
achieved

Sluggish pace of  
improvement in  
urban areas



**11 ministries supporting  
the program**

Sustainability of  
achievements not  
planned

Target shifted  
from 2020 to  
2018



***Mission Indradhanush: PM Modi calls  
for aggressive action plan to cover all  
children for immunization in a  
specific time-frame***



# Intensified Mission Indradhanush



**INTENSIFIED MISSION INDRADHANUSH LAUNCHED**

Ensuring full immunization to children and expecting mothers

- 1** To ensure full immunization in selected districts and cities to more than 90% by December 2018 against 2020 earlier
- 2** Four phases of the mission reached to 2.53 crore+ children and 68 lakh pregnant women
- 3** Progress to be monitored at the highest level under 'Proactive Governance and Timely Implementation' (PRAGATI)

The infographic features a vertical strip of four photographs on the left: a woman holding a child, a group of children holding certificates, a woman at a health center, and a group of women with children. At the bottom right is a small portrait of the Prime Minister. A rainbow logo with the text 'Get your child fully immunized' is in the top left corner.

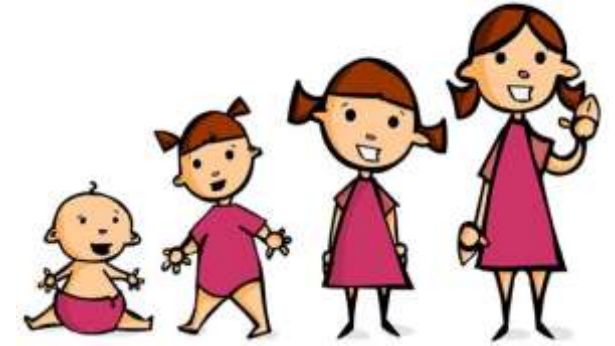
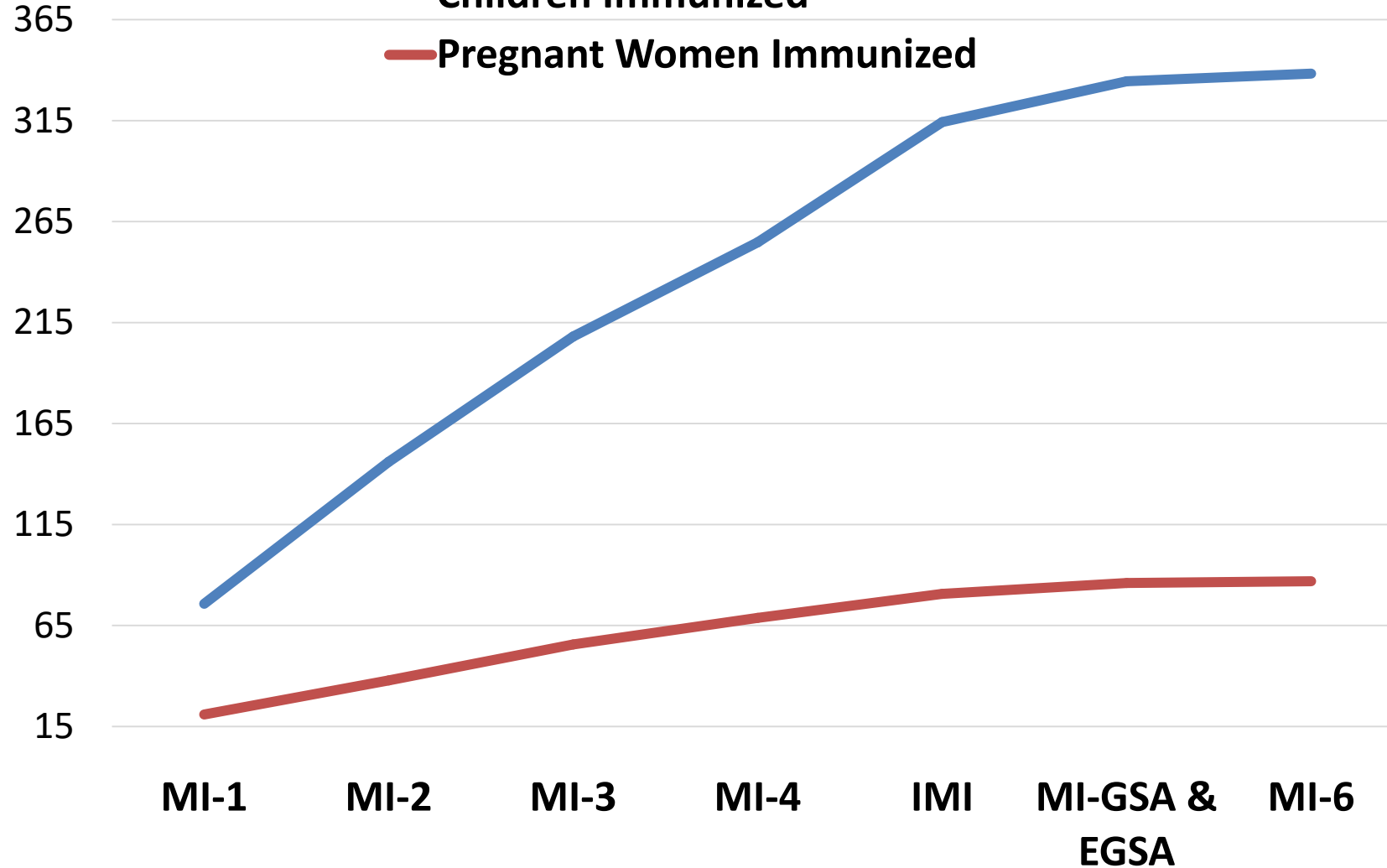


Hon'ble Prime Minister launched Intensified Mission Indradhanush on 8<sup>th</sup> October 2017

# Performance: Mission Indradhanush

Figures in lakh

— Children Immunized  
— Pregnant Women Immunized



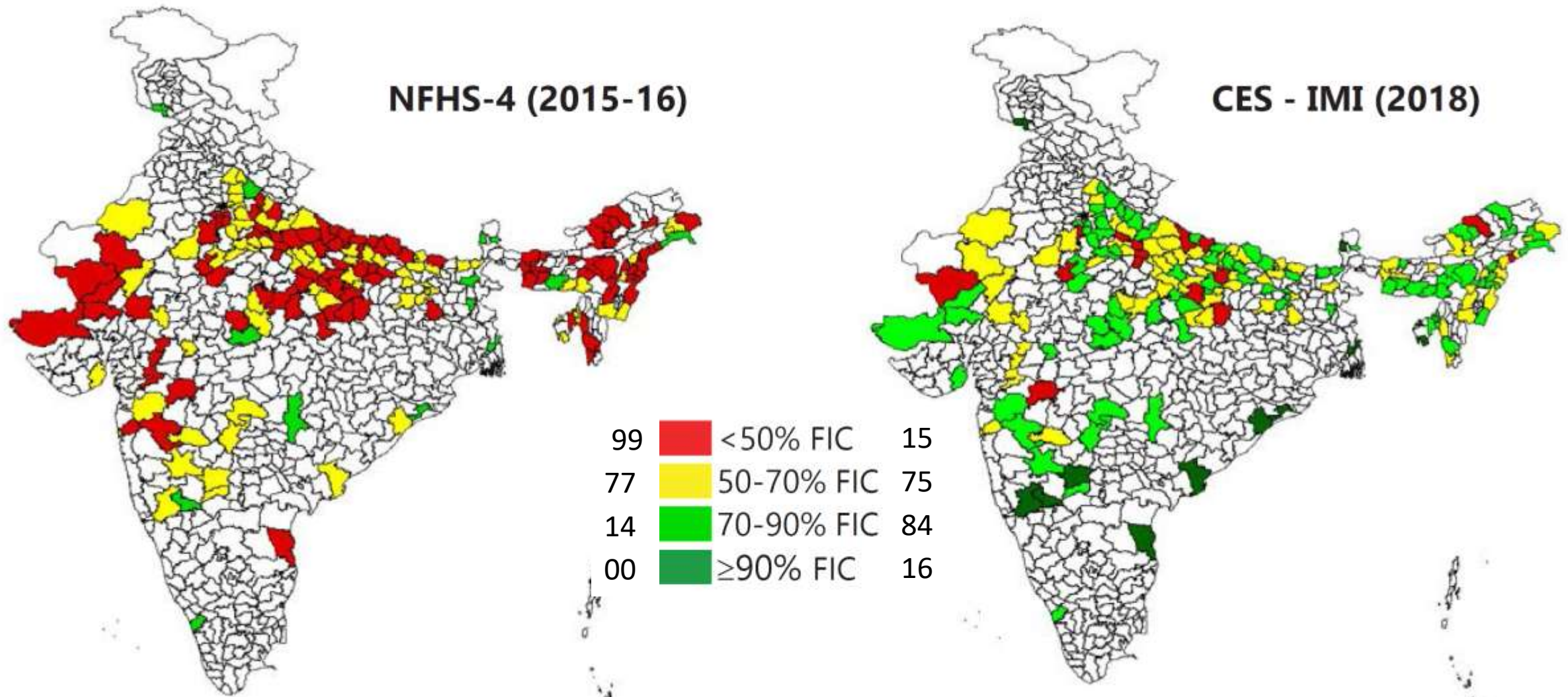
**3.39 crore children immunized**



**87.18 lakh  
pregnant  
women  
vaccinated**



# Impact of IMI in identified districts



***An average 18.5% increase in full immunization coverage as compared to NFHS-4 has been reported in 190 districts covered under IMI***



# MI under Gram Swaraj Abhiyan (GSA)/Extended GSA (EGSA)

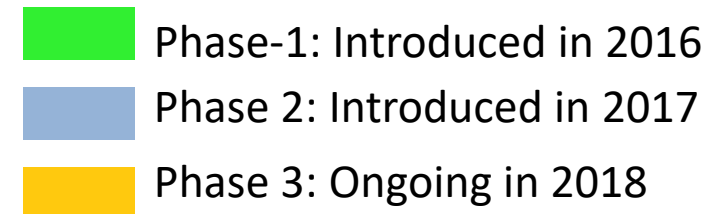
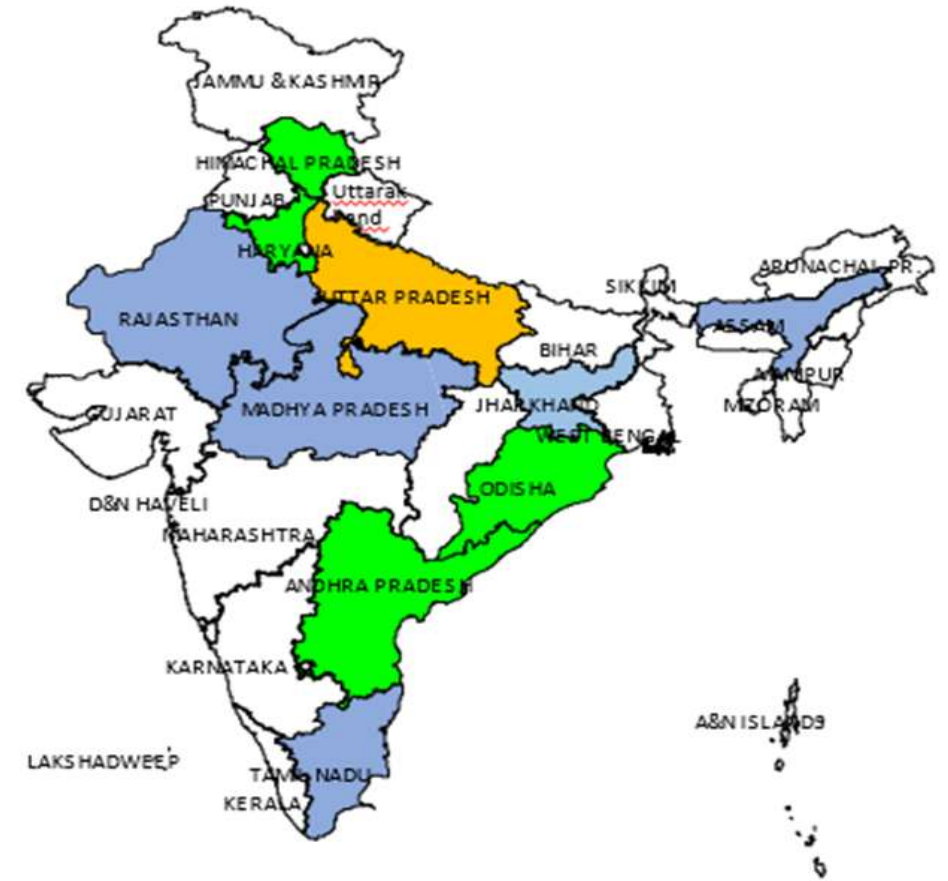
- MI under GSA - 16,850 villages across 25 states; and all UTs from Apr'18 to Jun'18
- MI – EGSA covered 48,929 villages across 117 aspirational districts. (7,408 villages in West Bengal did not participate).
- During MI in GSA/EGSA :
  - Children vaccinated: 20.22 lakh
  - Pregnant women vaccinated: 5.41 lakh



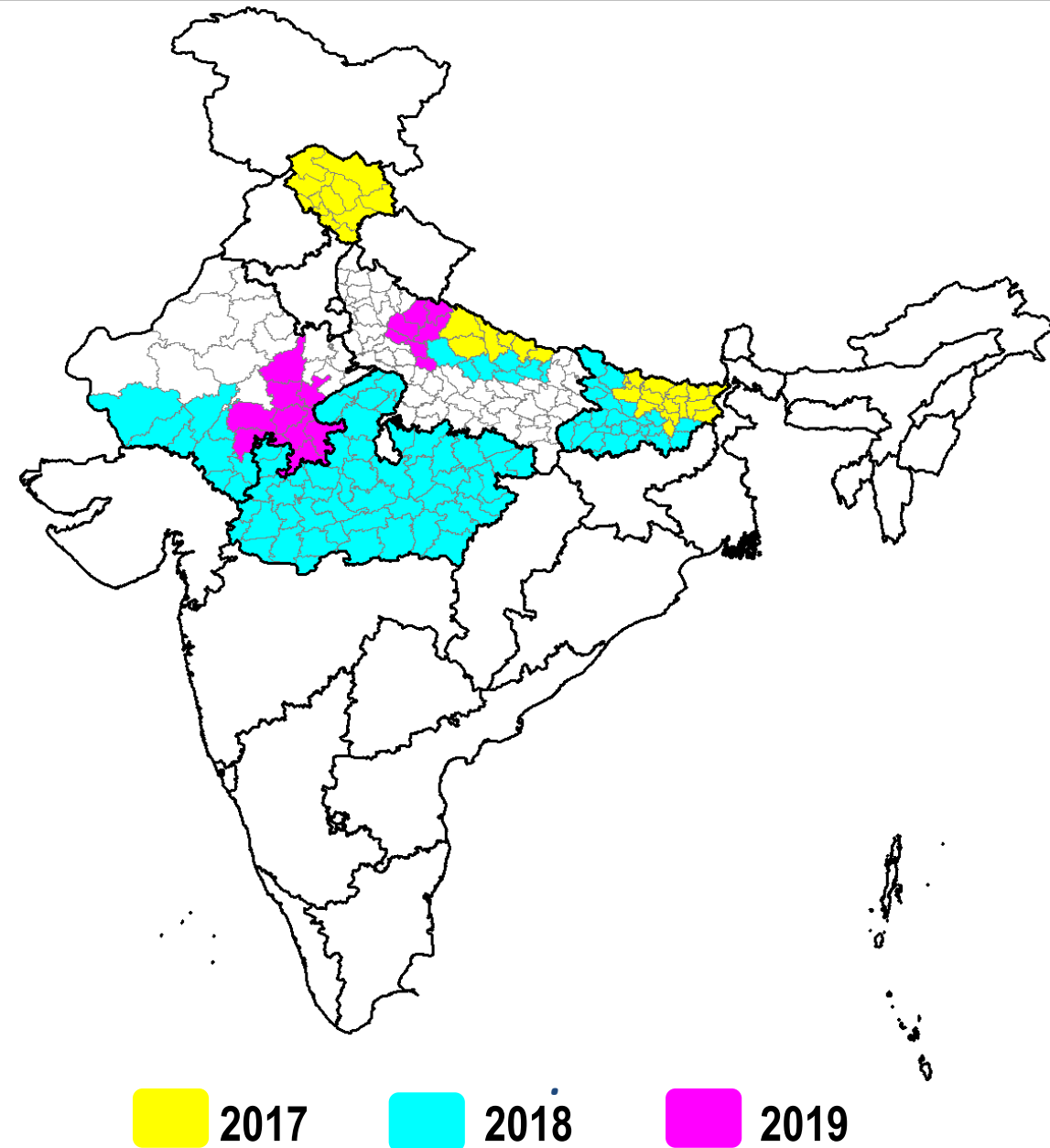
# New Vaccine Introduction

# Rotavirus vaccine Expansion Plan in India

- Criteria for State selection for RVV introduction
  - ✓ Diarrheal disease burden
  - ✓ AEFI preparedness
  - ✓ Routine immunization coverage and system preparedness
  - ✓ State willingness to introduce RVV
- Till March' 19, around 6.49 crore doses of Rotavirus vaccine have been administered to children.
- Expansion of Rotavirus vaccine under 'POSHAN Abhiyaan' to be done in all states in 2019-20 as per the directions of PMO



# Pneumococcal Conjugate Vaccine (PCV) Expansion Plan, India



- PCV has been introduced in Bihar, Himachal Pradesh, Madhya Pradesh, 19 districts of Uttar Pradesh and 18 districts of Rajasthan and Haryana (state initiative).
- Till March '19, around 116.89 lakh doses of PCV have been administered to children across above mentioned areas.
- In 2019, it will be further expanded to cover 9 and 7 additional districts in Rajasthan and Uttar Pradesh respectively.

## Percent birth cohort covered:

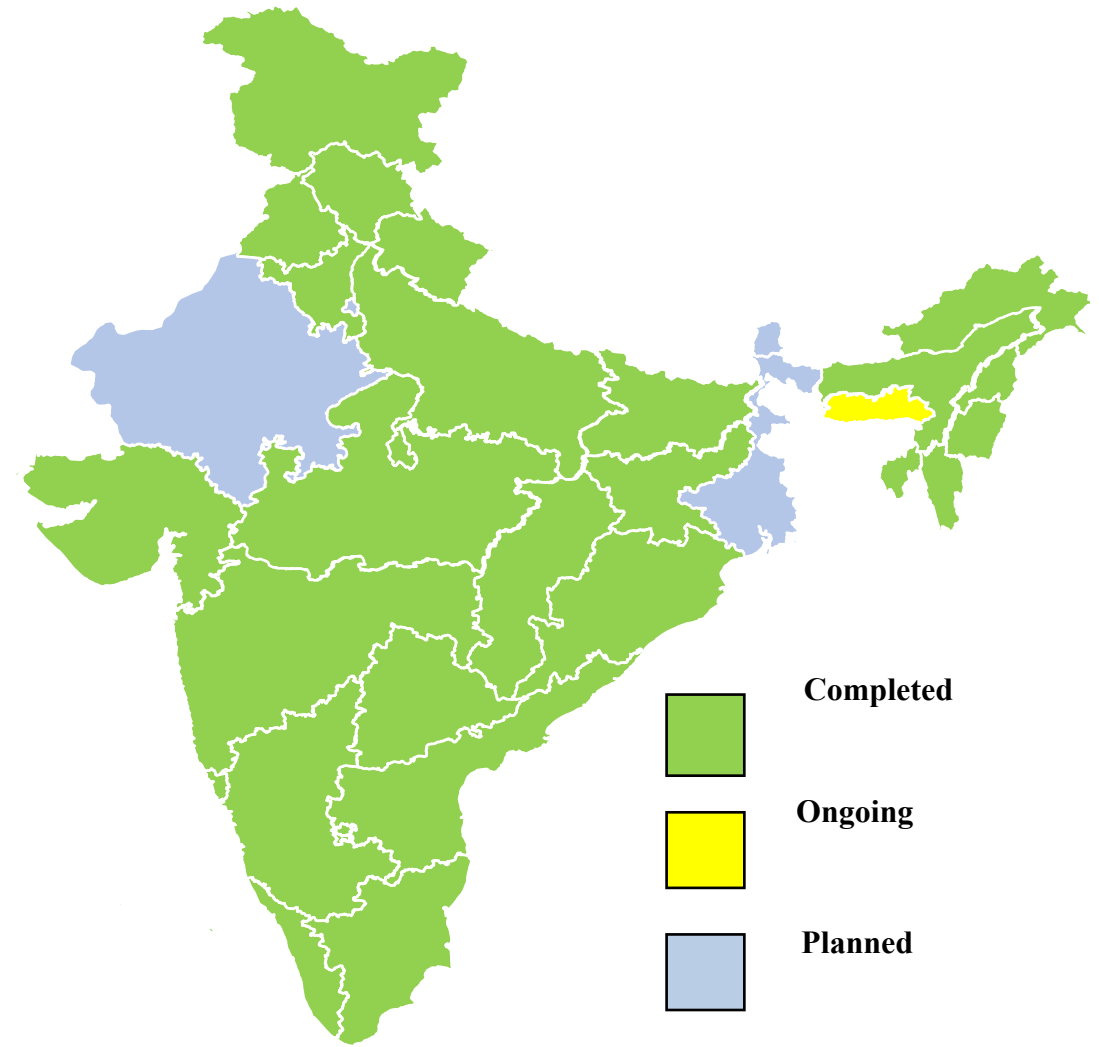
**Year-1 (2017):** Himachal Pradesh (100%), Bihar (50%),  
Uttar Pradesh (10%)

**Year-2 (2018):** Bihar (100%), Madhya Pradesh (100%),  
Rajasthan (25%) and Uttar Pradesh (20%)

**Year-3 (2019):** Rajasthan (50%) and Uttar Pradesh (30%)

# Measles Rubella (MR) Campaign

- WHO-SEARO goal of achieving Measles elimination by 2020, also reiterated by Hon'ble Finance Minister in the budget speech of 2017.
- Measles-Rubella vaccination campaign launched in Feb'17 targeting approx. 41 crore children aged 9 months-15 years across the country.
- Campaign has been completed in 31 states/UTs and ongoing in 1 state (Meghalaya).
- Subsequent to the completion of campaign, MR vaccine introduced in Routine Immunization replacing Measles vaccine at 9-12 months and 16-24 months of age.



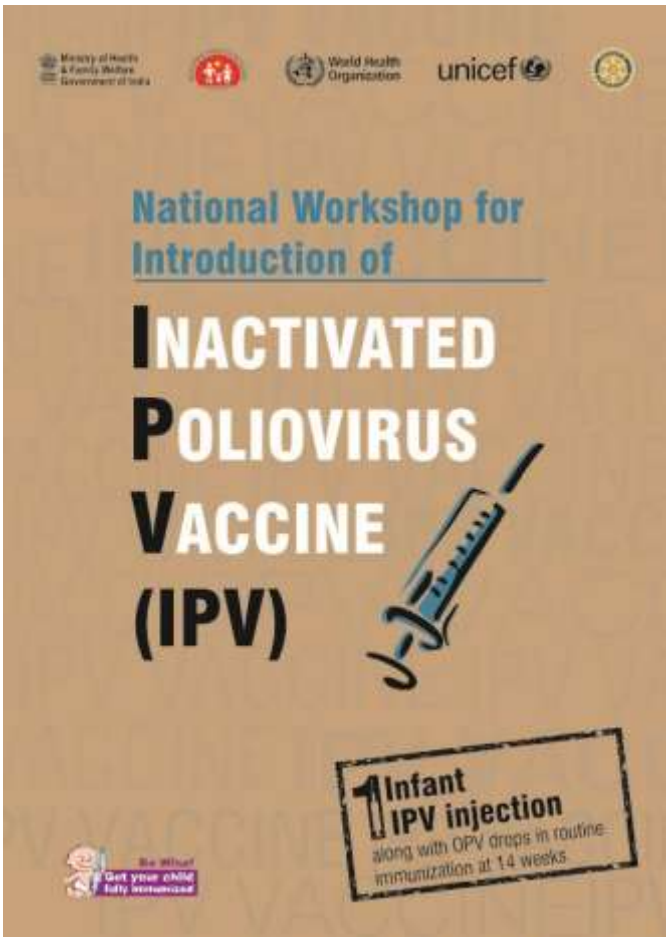
**>30.50 crore children vaccinated till date**



# MR Campaign Timelines – Remaining States

S. No	State/UT	Proposed Timeline
1.	West Bengal	-
2.	Rajasthan	July 2019
3.	Sikkim	August 2019
4.	Delhi	-

# Inactivated Polio Vaccine



- Launched on 30th November 2015, initially in 6 states
- Expanded to all states by April 2016
- 2 doses of fractional IPV (fIPV) given at 6 and 14 weeks of age of child
- Till March '19, around 8.89 crore doses of IPV vaccine have been administered to children across country

# Japanese Encephalitis(JE)

- JE vaccination: One time campaign strategy → single dose JE vaccine targeting **all** children from 1 to <15 years of age → JE vaccination is included into RI in endemic districts.
- **268 JE endemic districts** (including 37 identified in April'18) identified across **21 states** – campaigns completed in 230 districts → JE now part of RI.
- **Around 15.5 crore children** immunized during the campaign
- **35 high burden districts** (including 4 identified in April'18) identified in **3 states** for **Adult JE** vaccination in endemic blocks (Assam, UP, West Bengal).
- **Adult JE vaccination campaign** completed in 31 districts; **more than 3.3 crore beneficiaries** aged 15-65 years were vaccinated.



# Tetanus & adult Diphtheria (Td) vaccine

- Increase in immunization coverage in children led to shift in age-group of diphtheria cases to school going children and adults.
- Tetanus and adult Diphtheria (Td) vaccine has been recommended by National Technical Advisory Group on Immunization (NTAGI) in 2016.
- TT vaccine has been replaced by Td vaccine and will provide protection against both Tetanus and Diphtheria in adults.
- Td vaccine will replace 2 doses of TT or single booster dose of TT given to pregnant woman and booster doses at 10 and 16 years of age.

# Revised National Immunization Schedule

Age	Vaccines given
Birth	BCG, OPV-0, Hepatitis B Birth dose
6 Weeks	OPV-1, Pentavalent-1, fIPV-1, <b>Rota-1</b> & <b>PCV-1</b>
10 weeks	OPV-2, Pentavalent-2 & <b>Rota-2</b>
14 weeks	OPV-3, Pentavalent-3, fIPV-2, <b>Rota-3</b> & <b>PCV-2</b>
9-12 months	<b>MR-1</b> , JE1*, <b>PCV-Booster</b>
16-24 months	<b>MR-2</b> , JE2*, DPT-Booster 1, OPV- Booster
5-6 years	DPT-Booster 2
10 years	<b>Td</b>
16 years	<b>Td</b>
Pregnant Mother	<b>Td1, 2</b> or <b>Td Booster**</b>

\* in endemic districts only

\*\* one dose if previously vaccinated within 3 years

 Being introduced/scaled up

**National Immunization Schedule (NIS) for Infants, Children and Pregnant Women (Vaccine-wise)**

Vaccine	When to give	Dose	Route	Site
<b>For Pregnant Women</b>				
<b>Tetanus &amp; adult Diphtheria (Td)-1</b>	Early in pregnancy	0.5 ml	Intra-muscular	Upper Arm
<b>Td-2</b>	4 weeks after Td-1	0.5 ml	Intra-muscular	Upper Arm
<b>Td- Booster</b>	If received 2 TT/Td doses in a pregnancy within the last 3 years*	0.5 ml	Intra-muscular	Upper Arm



<b>For Infants</b>				
<b>Bacillus Calmette Guerin (BCG)</b>	At birth or as early as possible till one year of age	0.1ml (0.05ml until 1 month age)	Intra-dermal	Left Upper Arm
<b>Hepatitis B - Birth dose</b>	At birth or as early as possible within 24 hours	0.5 ml	Intra-muscular	Antero-lateral side of mid-thigh
<b>Oral Polio Vaccine (OPV)-0</b>	At birth or as early as possible within the first 15 days	2 drops	Oral	Oral
<b>OPV 1, 2 &amp; 3</b>	At 6 weeks, 10 weeks & 14 weeks (OPV can be given till 5 years of age)	2 drops	Oral	Oral
<b>Pentavalent 1, 2 &amp; 3</b>	At 6 weeks, 10 weeks & 14 weeks (can be given till one year of age)	0.5 ml	Intra-muscular	Antero-lateral side of mid-thigh
<b>Pneumococcal Conjugate Vaccine(PCV)</b>	Two primary doses at 6 and 14 weeks followed by Booster dose at 9-12 months	0.5 ml	Intra-muscular	Antero-lateral side of mid-thigh
<b>Rotavirus (RVV)</b>	At 6 weeks, 10 weeks & 14 weeks (can be given till one year of age)	5 drops (liquid vaccine) 2.5 ml (lyophilized vaccine)	Oral	Oral

<b>Inactivated Polio Vaccine (IPV)</b>	Two fractional dose at 6 and 14 weeks of age	0.1 ml	Intra dermal two fractional dose	Intra-dermal: Right upper arm
<b>Measles Rubella (MR) 1<sup>st</sup> dose</b>	9 completed months-12 months. (Measles can be given till 5 years of age)	0.5 ml	Sub-cutaneous	Right upper Arm
<b>Japanese Encephalitis (JE) - 1</b>	9 completed months-12 months.	0.5 ml	Sub-cutaneous (Live attenuated vaccine)  Intramuscular(Killed vaccine)	Left upper Arm (Live attenuated vaccine)  Anterolateral aspect of mid thigh (Killed vaccine)
<b>Vitamin A (1<sup>st</sup> dose)</b>	At 9 completed months with measles-Rubella	1 ml ( 1 lakh IU)	Oral	Oral

<b>For Children</b>				
<b>Diphtheria, Pertussis &amp; Tetanus (DPT) booster-1</b>	16-24 months	0.5 ml	Intra-muscular	Antero-lateral side of mid-thigh
<b>MR 2<sup>nd</sup> dose</b>	16-24 months	0.5 ml	Sub-cutaneous	Right upper Arm
<b>OPV Booster</b>	16-24 months	2 drops	Oral	Oral
<b>JE-2</b>	16-24 months	0.5 ml	Sub-cutaneous (Live attenuated vaccine)  Intramuscular(Killed vaccine)	Left upper Arm (Live attenuated vaccine)  Anterolateral aspect of mid thigh (Killed vaccine)
<b>Vitamin A (2<sup>nd</sup> to 9<sup>th</sup> dose)</b>	16-18 months. Then one dose every 6 months up to the age of 5 years.	2 ml (2 lakh IU)	Oral	Oral
<b>DPT Booster-2</b>	5-6 years	0.5 ml.	Intra-muscular	Upper Arm
<b>Td</b>	10 years & 16 years	0.5 ml	Intra-muscular	Upper Arm

\*One dose if previously vaccinated within 3 years

Note:

- JE Vaccine is introduced in select endemic districts after the campaign.
- The 2<sup>nd</sup> to 9<sup>th</sup> doses of Vitamin A can be administered to children 1-5 years old during biannual rounds, in collaboration with ICDS.
- PCV in selected states/districts: Bihar, Himachal Pradesh, Madhya Pradesh, Rajasthan & Uttar Pradesh (selected districts), and in Haryana as state initiative



# Improving Quality

# Vaccine Logistics & Cold Chain Management

- National Cold Chain Resource Centre (NCCRC), Pune and National Cold Chain & Vaccine Management Resource Centre (NCCVMRC) -NIHFW, New Delhi established to provide technical training to cold chain technicians in repair & maintenance of cold chain equipment.
- National Cold Chain Management Information System (NCCMIS) to track cold chain equipment inventory, availability and functionality.

# National Effective Vaccine Management (EVM) Assessment 2018

Diagnostic tool to assess and review three “P”s - Process, Practices and Policies of Efficient Immunization Supply Chain-Cold Chain – Supported by comprehensive Improvement plan

Participation by -  
MoHFW, Medical  
Colleges (16), ITSU,  
NCCVMRC, UNICEF,  
UNDP, WHO, JSI.

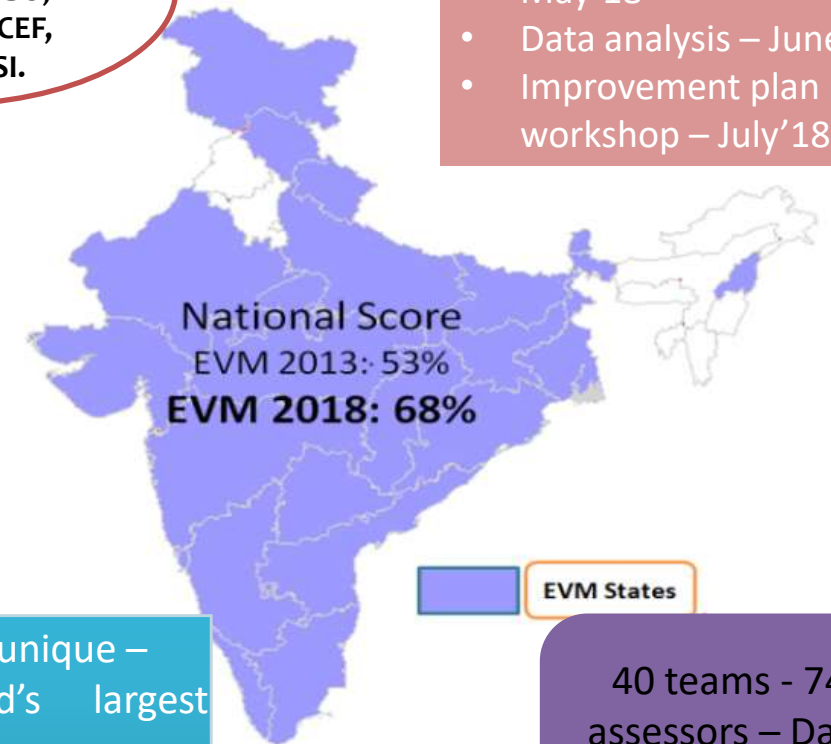
Status –

- Data collection - May'18
- Data analysis – June'18
- Improvement plan workshop – July'18

## *Under 9 Global Criteria's*

1. Vaccine Arrival Process
2. Vaccine Storage Temperature
3. Storage Capacity
4. Building, CCE & Transport
5. Maintenance & Repair
6. Stock Management
7. Distribution
8. Vaccine Management Practices
9. MIS & Supportive Functions

2018 – 23 states



What make this assessment unique –

- EVM 2018 is world's largest assessment
- Participation by players from different domains of public health
- Mobile Based Paper less assessment
- Shortest duration (2 months) – EVM Assessment

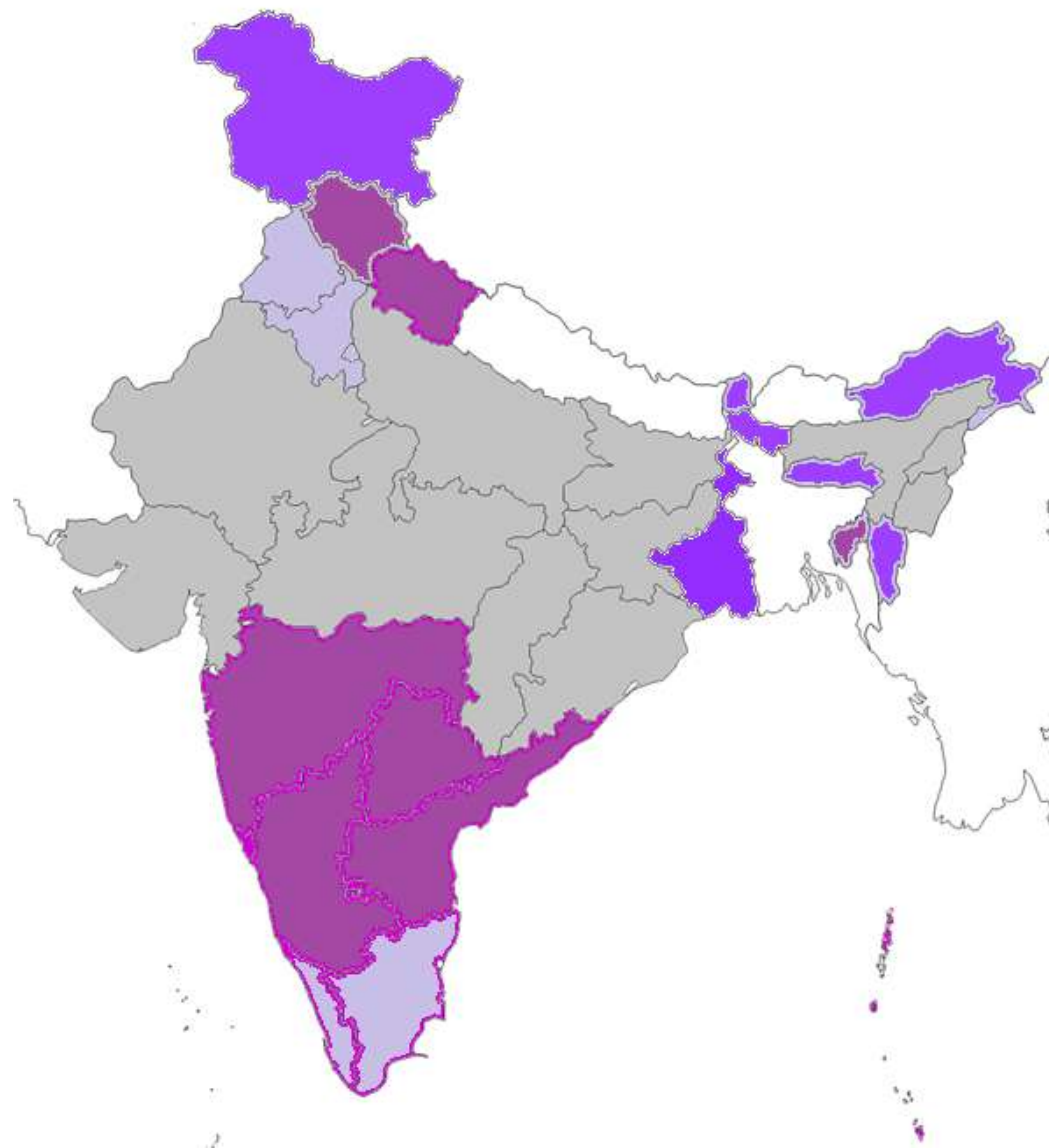
40 teams - 74  
assessors – Data  
collection from  
145 sites



# eVIN status and scale up plan

Electronic Vaccine Intelligence Network (eVIN) rollout for :

Real time stock management and  
Real time monitoring of cold chain temperature using mobile technology and data logger (sim based)



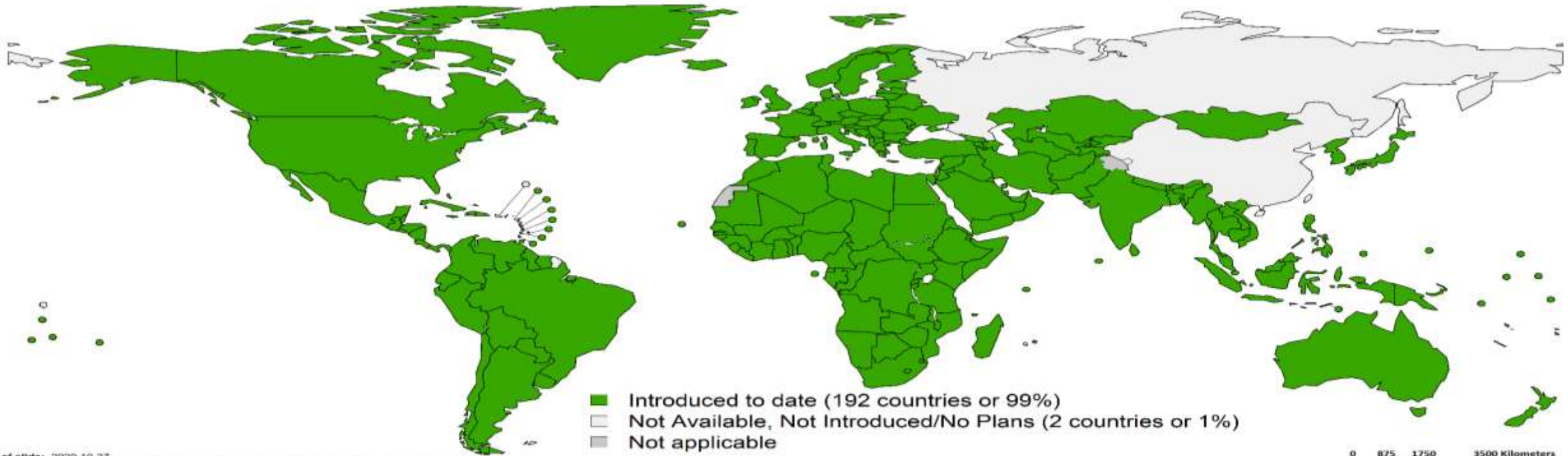
**Phase 4, Initiation  
planned in July 2019**

**Phase 3 , Initiation  
planned in October 2018**

**Phase 2 Implementation  
initiated. Expected  
completion by June 2019**

**Current eVIN States**

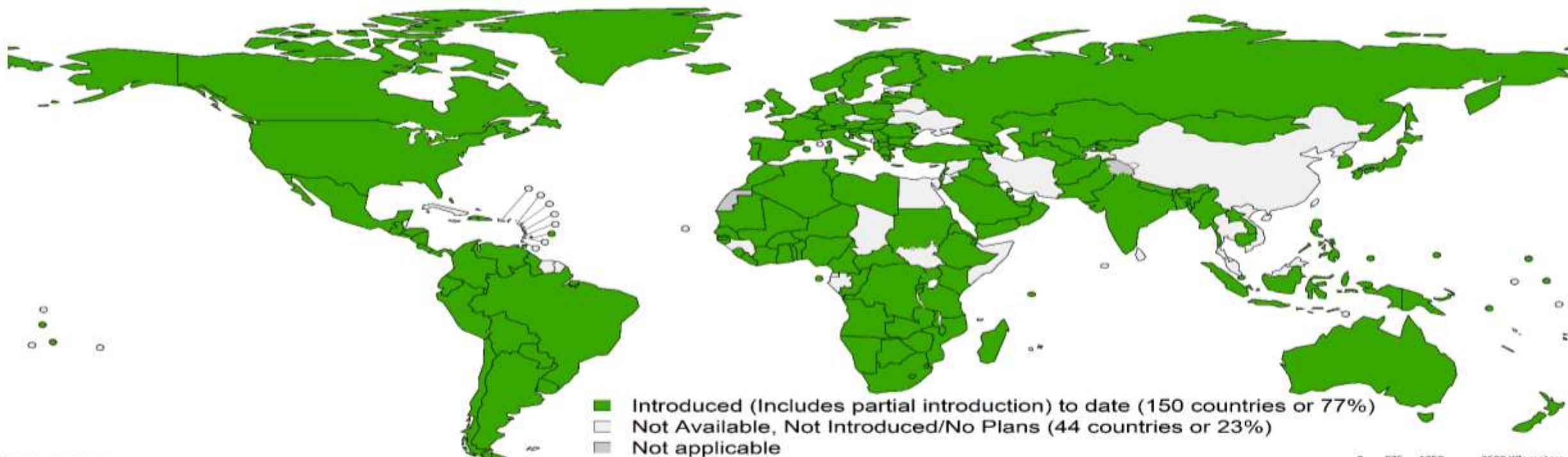
# Countries with Hib vaccine in the national immunization programme



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Data source: IVB database as at 23th October 2020

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# Countries with Pneumococcal Conjugate vaccine in the national immunization programme



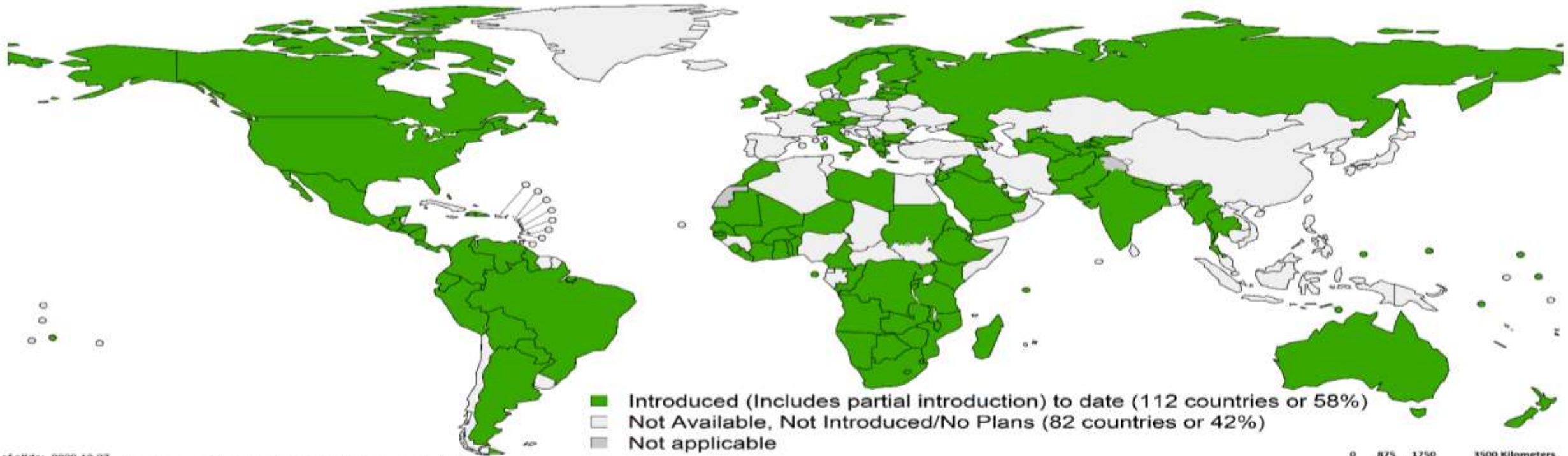
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# Countries with Rotavirus vaccine in the national immunization programme



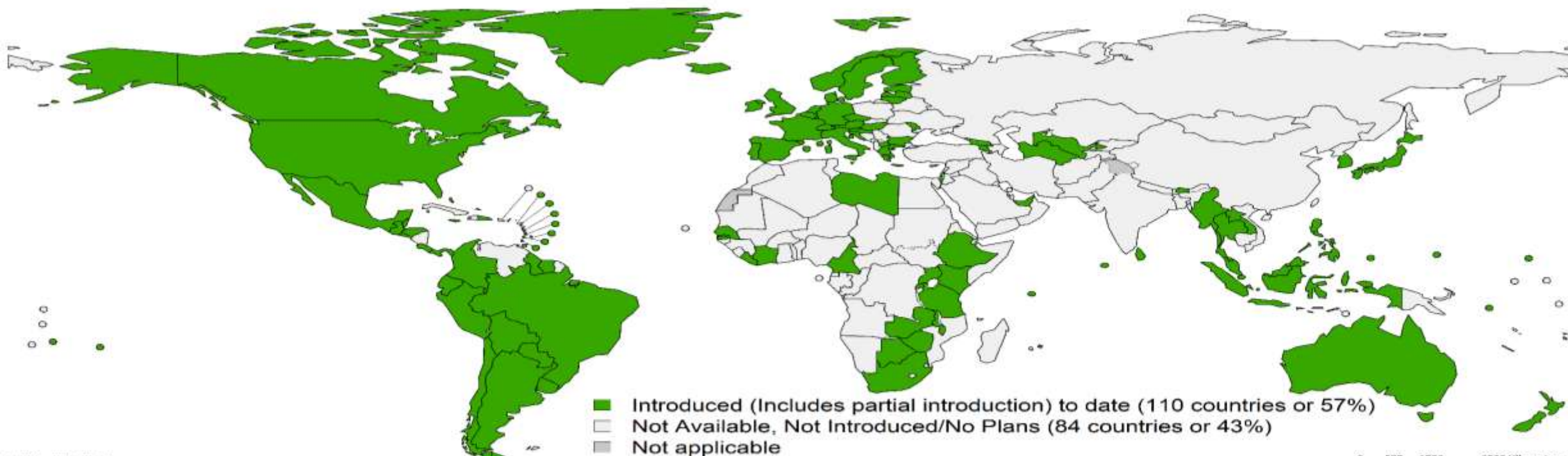
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0 875 1750 3500 Kilometers



# Countries with HPV vaccine in the national immunization programme

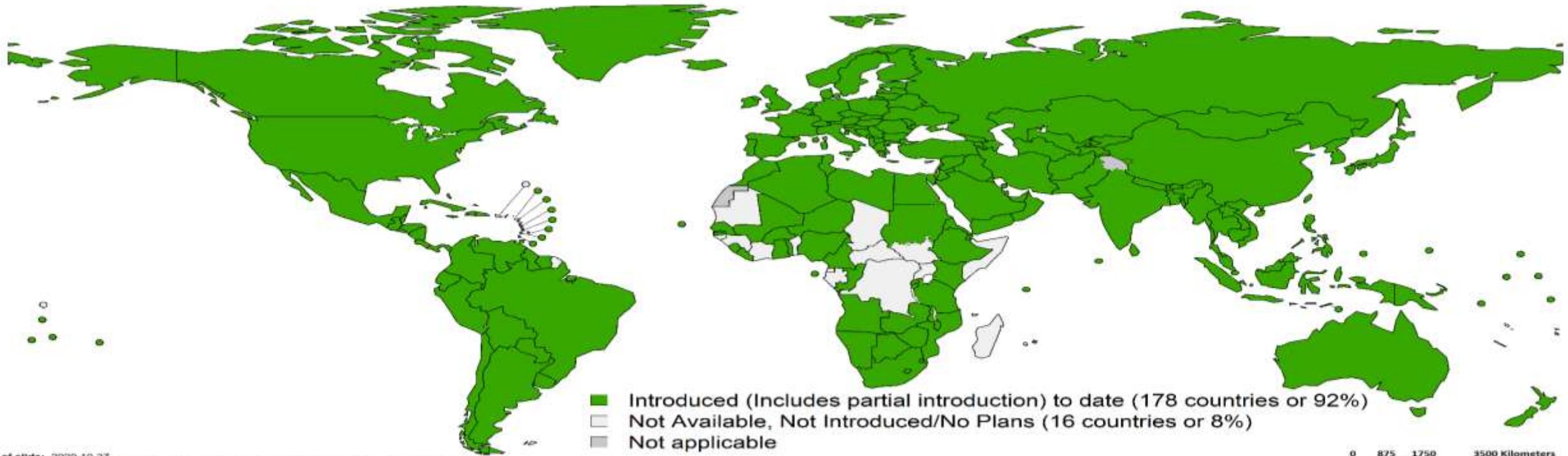


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# Countries with Measles second dose vaccine in the national immunization programme



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# Countries with Rubella containing vaccine in the national immunization programme

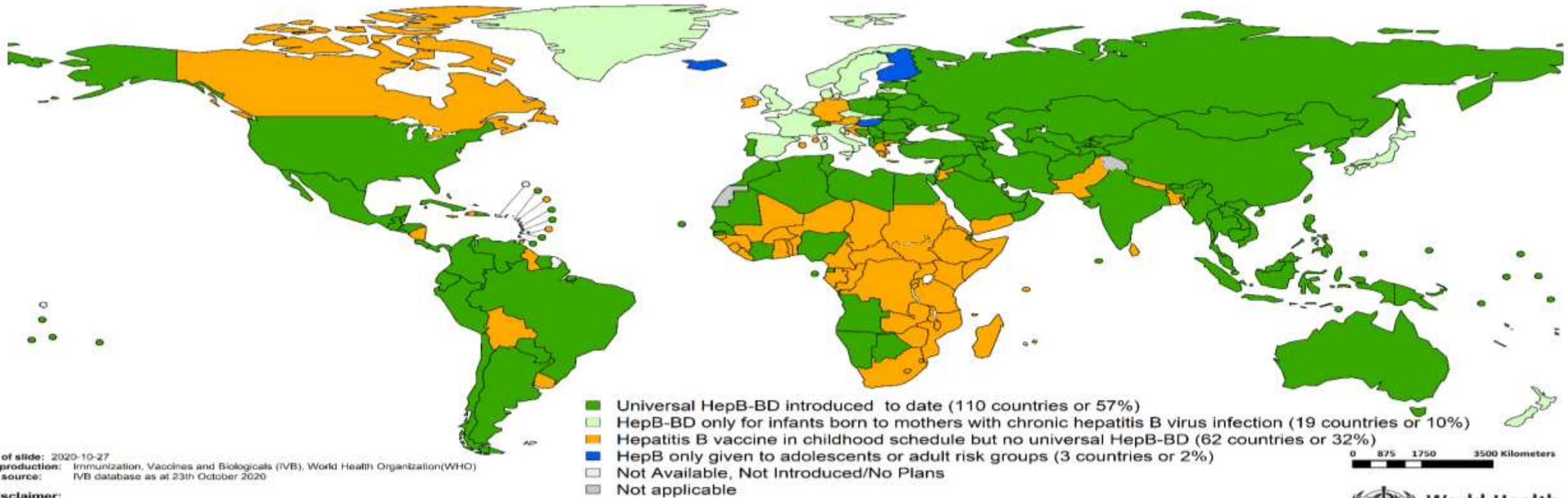


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# Hepatitis B Birth dose (HepB-BD) vaccination strategies in the national immunization programme



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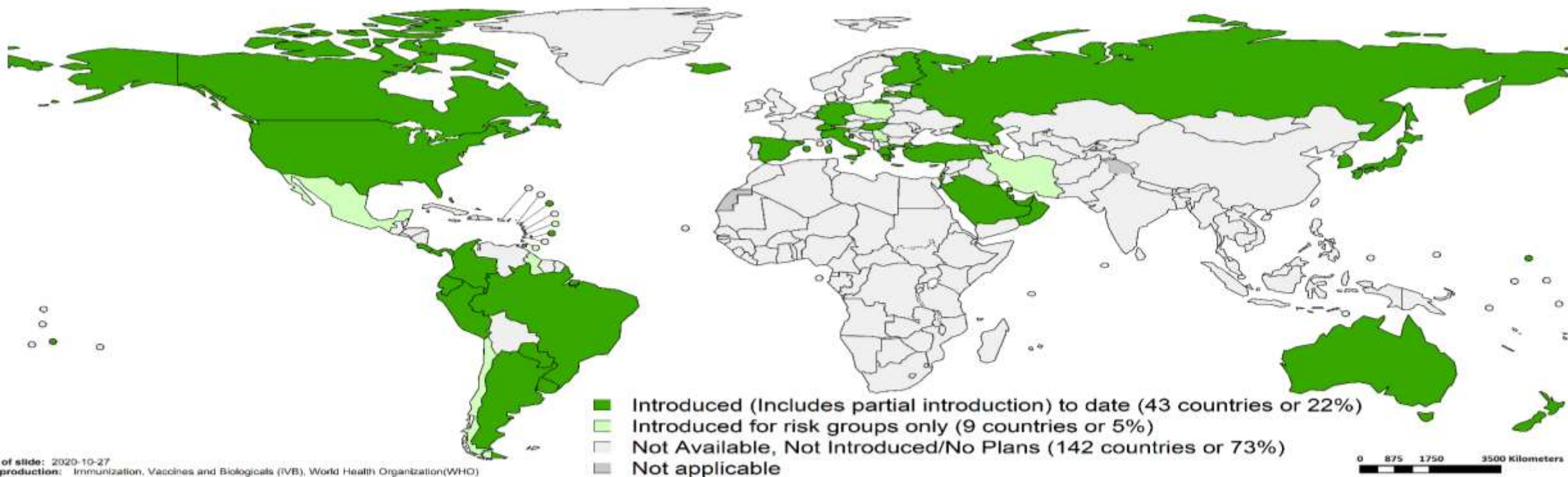
# Countries with Mumps containing vaccine in the national immunization programme



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# Countries with Varicella containing vaccine in the national immunization programme



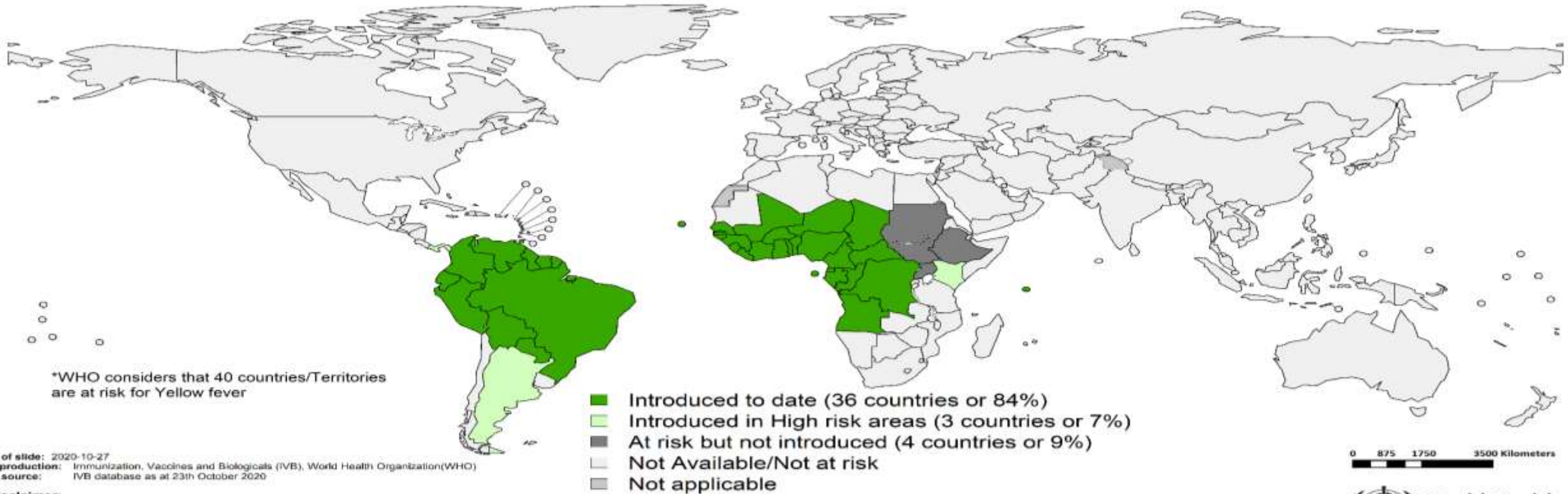
Date of slide: 2020-10-27  
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# Countries and Territories using Yellow fever\* vaccine to date

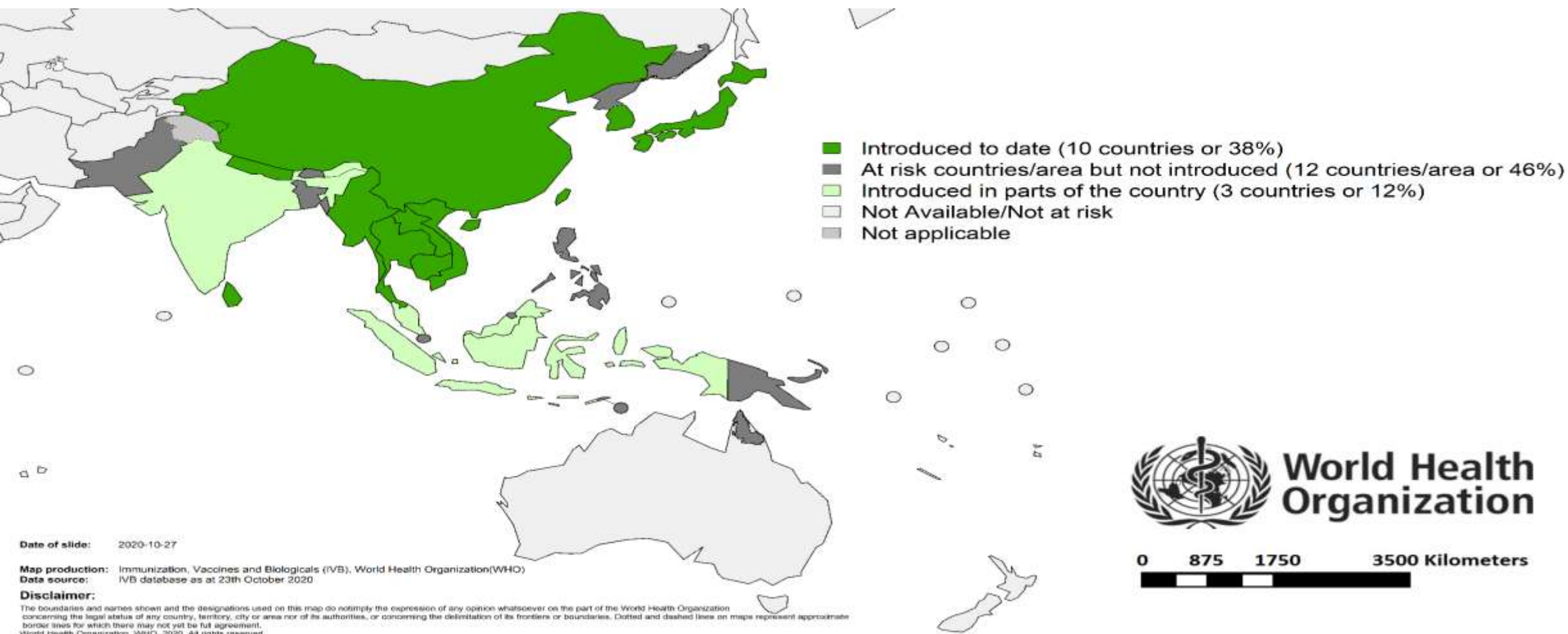


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# Countries and Territories using JE vaccine to date



# Do's and Don'ts during immunization sessions

Do's	Don'ts
<b>Vaccination schedule</b>	
<ul style="list-style-type: none"> <li>It is safe and effective to give BCG, DPT, OPV and Measles vaccines at the same time to a child who has completed 9 months and never been vaccinated.</li> <li>Give BCG to infants less than 1 year of age (never give BCG to children above 1 year of age).</li> <li>If a child is brought late for a dose, pick up where the schedule was left off. For example, if a child left with DPT-2 and comes after 3 months give DPT-3.</li> </ul>	<ul style="list-style-type: none"> <li>Withhold the vaccine in case of illness such as cold, cough, diarrhoea or fever.</li> </ul>
<b>Cold chain</b>	
<ul style="list-style-type: none"> <li>Check expiry date and VVM label of vaccine vial before immunizing every child.</li> <li>Keep the vaccines and diluents in a plastic bag/zipper bag in the centre of vaccine carrier with 4 conditioned ice-packs. Make sure that the diluents are also at +2 to +8 centigrade before reconstitution.</li> <li>Take one ice pack from vaccine carrier and keep reconstituted BCG &amp; Measles vaccines only on the top of the ice pack.</li> </ul>	<ul style="list-style-type: none"> <li>Leave vaccine carrier in sunlight: this spoils vaccines that are sensitive to heat and light.</li> <li>Leave the lid open, this can allow heat and light into the carrier, which can spoil vaccine.</li> <li>Drop or sit on the vaccine carrier: this can damage the carrier.</li> <li>Carry vaccines in handbag as this can spoil vaccines that are sensitive to heat.</li> <li>Keep the DPT, DT, TT and Hep. B vaccines on the ice pack during the session.</li> </ul>
<b>Vaccine handling and administration</b>	
<ul style="list-style-type: none"> <li>Welcome beneficiaries.</li> <li>Wash hands before conducting the session.</li> <li>Verify beneficiary's record and age of the child.</li> <li>Screen for contraindications.</li> <li>Check label of the vial and expiry date.</li> <li>Lightly shake the vial of T-Series Vaccine before drawing the dose.</li> <li>Use a new AD syringe for each injection and new disposable syringe for each reconstitution.</li> <li>Use correct diluent for reconstitution of vaccine.</li> <li>Give appropriate vaccine.</li> <li>Inject vaccine using the correct site and route for the vaccine e.g. Intradermal in left arm for BCG; subcutaneous in right arm for Measles; intramuscular in anterolateral aspect of mid thigh for DPT and Hepatitis B.</li> <li>Allow dose to self-disperse instead of massaging.</li> <li>Explain potential adverse events following immunization and what to do.</li> <li>Discuss with beneficiaries/parents about next visit.</li> </ul>	<ul style="list-style-type: none"> <li>Use un-sterile syringe or needle for immunization.</li> <li>Draw air into AD syringes.</li> <li>Touch any part of the needle.</li> <li>Recap the needle.</li> <li>Leave the needle inside the vial.</li> <li>Ever inject in the buttock, never do that.</li> <li>Massage the vaccination site after vaccination.</li> <li>Use reconstituted measles and BCG vaccine after 4hrs and JE after 2 hrs.</li> <li>Use vaccine with VVM in unusable stage or with expiry date.</li> </ul>
<b>Recording and reporting</b>	
<ul style="list-style-type: none"> <li>Fully document each immunization in the immunization card, tally sheet and immunization register. Ask parents/guardians to bring the card on next visit.</li> <li>Retain the counterfoil.</li> </ul>	<ul style="list-style-type: none"> <li>Turn away beneficiaries for not bringing the card.</li> <li>Leave any cell blank in immunization card.</li> </ul>
<b>Adverse events following immunization (AEFI)</b>	
<ul style="list-style-type: none"> <li>In case of serious AEFI refer the patient to appropriate health facility, inform your supervisor immediately – document the type of vaccine(s), batch number, expiry date, and full address of the child.</li> <li>Report all serious AEFIs to the MOI/C.</li> </ul>	<ul style="list-style-type: none"> <li>Report minor reaction following vaccination (mild fever of less than three days, redness and pain).</li> </ul>
<b>Social mobilization</b>	
<ul style="list-style-type: none"> <li>Use vaccination card to remind parents when to return with their child.</li> <li>Enlist community team like AWW, ASHA, NGOs and other community-based workers to remind parents of the importance of full immunization.</li> </ul>	<ul style="list-style-type: none"> <li>Leave any community meeting without communicating about immunization session days.</li> </ul>

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- <http://www.nihfw.org/doc/NCHRC-Publications/>
- [https://www.who.int/immunization/monitoring surveillance/VaccineIntroStatus/](https://www.who.int/immunization/monitoring_surveillance/VaccineIntroStatus/)
- [Universal Immunisation Programme | National Health Portal Of India \(nhp.gov.in\)](#)
- Park's Textbook of Preventive and Social Medicine, 25<sup>th</sup> Edition







<u>Author</u>	<u>Study</u>	<u>Method</u>	<u>Result</u>	<u>Conclusion</u>
<a href="#">Latika Nath, Prabhdeep Kaur,<sup>1</sup> and Saurabh Tripathi<sup>2</sup></a> <a href="#">Indian J Community Med.</a> 2015 Oct-Dec; 40(4): 239–245	cross-sectional survey (low level of evidence)	A facility, session site and cross-sectional survey of 180 children were done and proportions for various indicators were estimated. We determined factors associated with not taking vaccination using multivariate analysis.	The highest prevalent districts in Karnataka are Dakshina Kannada for malaria, Udupi for dengue, Tumakuru for chikungunya and Bidar for lymphatic filariasis. Major factors related to poor outcome in some districts were rigid attitude of the community, poor support of local panchayats and less human resources.	There was low immunization coverage among migrants within adequate supervision, poor cold chain maintenance, and improper tracking of dropouts. Mobile immunization teams, prelisting of migrant children, and change in incentives of ASHAs for child tracking were needed. A monitoring plan for sessions and cold chain needed enforcement.