

# Current Strategies for Vaccination Against Hepatitis A

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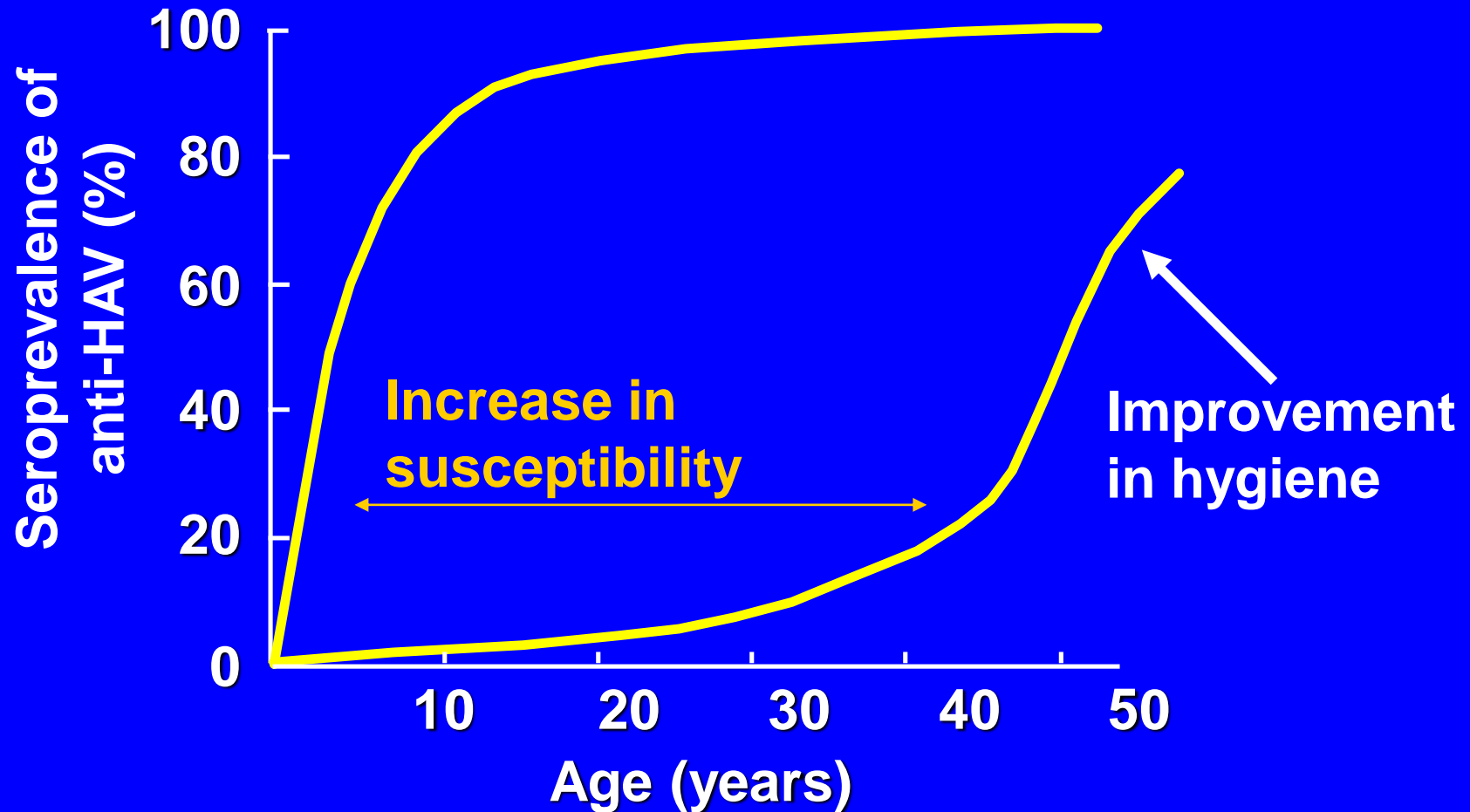
# Disclosure - Hepatitis A

- Advisor -Viral Hepatitis Prevention Board
- Advisor- WHO
- Advisor-European Centers for Disease Control

# Outline

- Epidemiology
- Hepatitis A vaccines
- Control and prevention strategies
- Global overview of hepatitis A vaccination programs
- Elimination of HAV infection in Israel
- Single dose immunization-Argentina
- Safety
- Summary

# Hepatitis A epidemiology shifts with improving hygiene

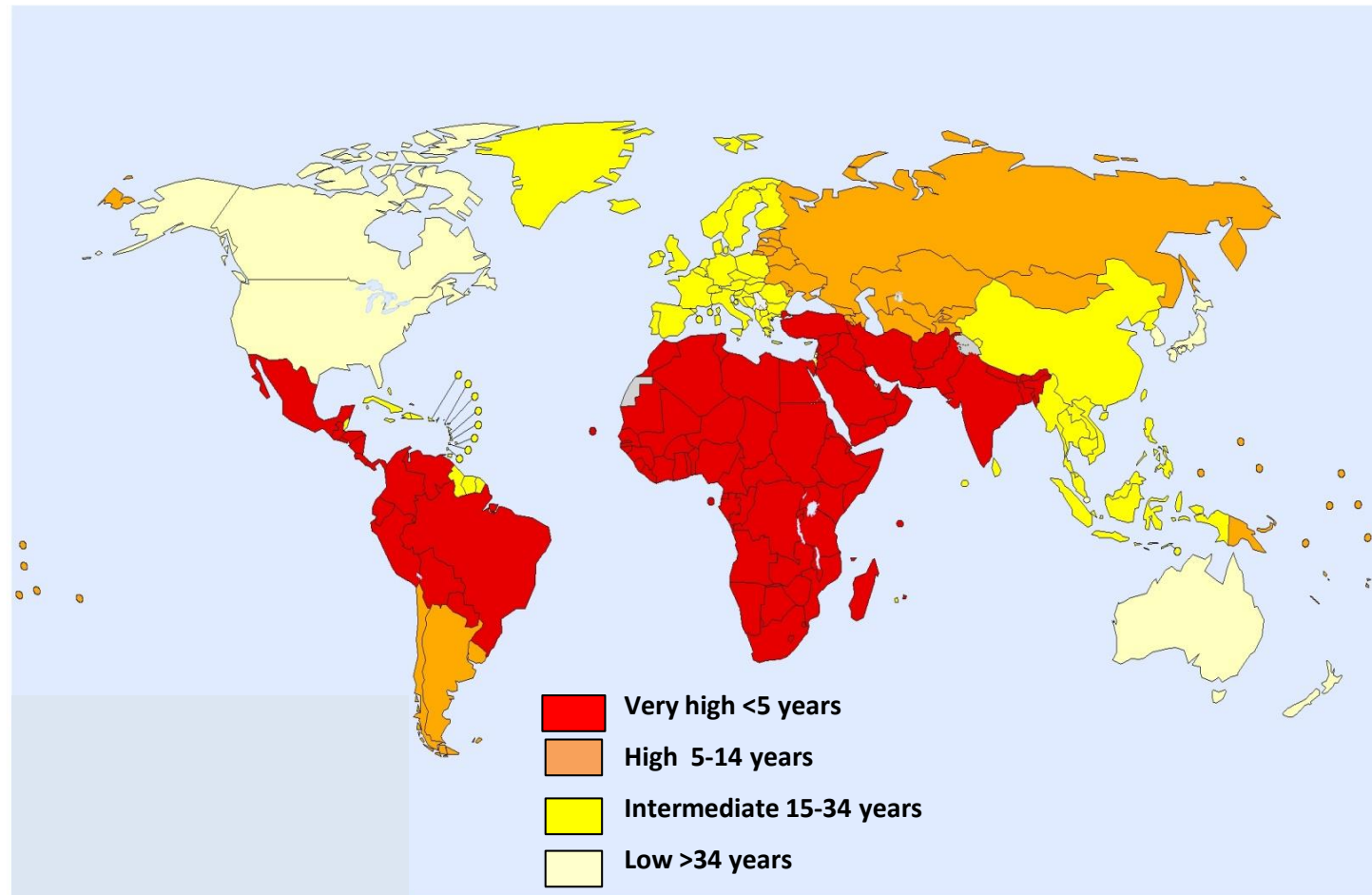


# WHO position paper on epidemiology of hepatitis A vaccines – June 2012\*

Based on an ongoing reassessment of the global burden of hepatitis A, WHO estimates suggest :

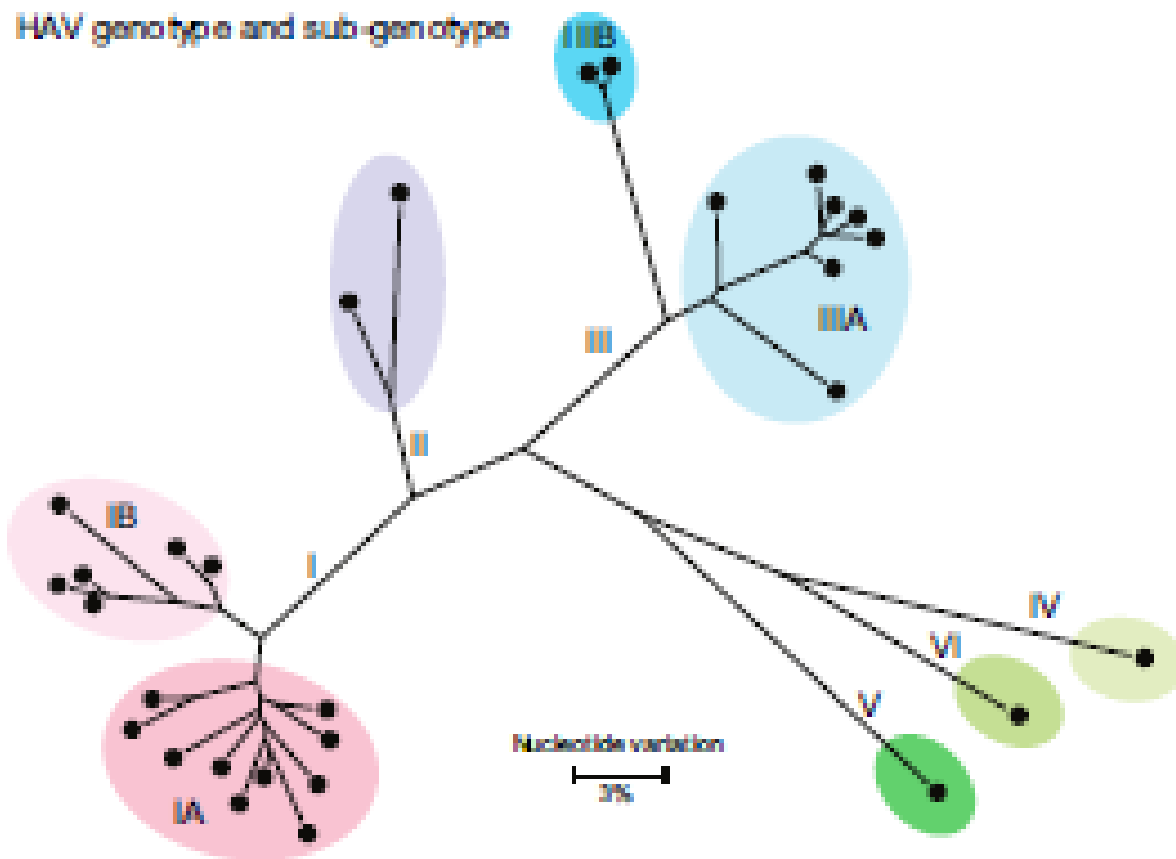
- An increase in the number of acute hepatitis A cases from 177 million in 1990 to 212 million in 2005
- Deaths due to hepatitis A to increase from 30 283 in 1990 to 35 245 in 2005.
- Increased numbers of cases estimated to occur in the age groups 2–14 years and >30 years
- Adult infections
  - 75-90% of cases are symptomatic
  - Historically- fulminant hepatitis is rare ( $\leq 1\%$ ) but rising incidence of fulminant hepatitis in distinct regions?
  - 1.75-2.1% mortality rate after  $\geq 40$  years of age

# Global risk map of HAV immunity in 2005: Age at midpoint of population immunity to HAV\*



\*Mohd Hanafiah K, Jacobsen KH, Wiersma ST. Challenges to mapping the health risk of hepatitis A virus infection. *Int J Health Geogr* 2011;10:57

Jacobsen KH, Wiersma ST. Hepatitis A virus seroprevalence by age and world region, 1990 and 2005. *Vaccine* 2010;28:6653-



**Fig. 6. HAV genotype classification.** Phylogenetic analysis of the six currently recognised HAV genotypes. Reproduced with permission from.<sup>72</sup> HAV, hepatitis A virus.

# Milestones in development of hepatitis A vaccines\*

- **1988** - Propagation of attenuated HAV in culture
- **Early 1990s** - Pivotal efficacy studies in Thailand and the US
- **1996** - ACIP-Introduction of vaccine to selected risk groups
- **1999** >- Universal vaccination in selected regions/countries
- **2005** - Single dose immunization
- **2007** - Post exposure prophylaxis
- **2007** - Int. meeting: Global control of HAV infection, Miami \*



# Hepatitis A vaccines

- Inactivated Vaccines

Vs

- Live, attenuated vaccine
- 

- Monovalent Vaccines

Vs

- Combined vaccines

# Hepatitis A vaccines

## Inactivated, Monovalent or Combined vaccines

- ✓ Manufactured from attenuated HAV strain
- ✓ Formaldehyde, inactivated
- ✓ Contain an adjuvant:
  - aluminum hydroxide
  - or
  - formulated in virosomes

## ❖ Live attenuated vaccines

No adjuvant

## Inactivated Vs live attenuated vaccines

Differences in:

- ➔ Technology of production
- ➔ Cost
- ➔ Pace of immune response to vaccination
- ➔ Surveillance of safety and tolerability
- ➔ Distribution

# Inactivated Vs Live Attenuated HAV Vaccines

**Table 1. Monovalent formalin-inactivated hepatitis A vaccines.\*#**

Attenuated HAV strain	Trade name	Adjuvant	HAV antigen Dose/injection		Manufacturers
			Paediatric	Adult	
HM-175	HAVRIX®	Aluminium hydroxide	720 EU	1440 EU	GSK
CR-326	VAQTA®	Aluminium hydroxide	25 U	50 U	MSD
GBM	AVAXIM®	Aluminium hydroxide	80 U	160 U	Aventis Pasteur
TZ84	HEALIVE®	Aluminium hydroxide	250 U	500U	Sinovac Biotech Co LTd
Lv-8	Weisairuian®	Aluminium hydroxide	320 EU	640 EU	Inst Med Biol
YN5	Veraxim®	Aluminium hydroxide	800 EU	1600 EU	Shanghai Wison Bioengineering Inc
RG-SB	EPAXAL®	Virosomes	24 U	24 U	Crucell/Berna Biotech

\* Modified and updated from references <sup>75,160</sup>

**Table 2. Live attenuated hepatitis A vaccines.\*\***

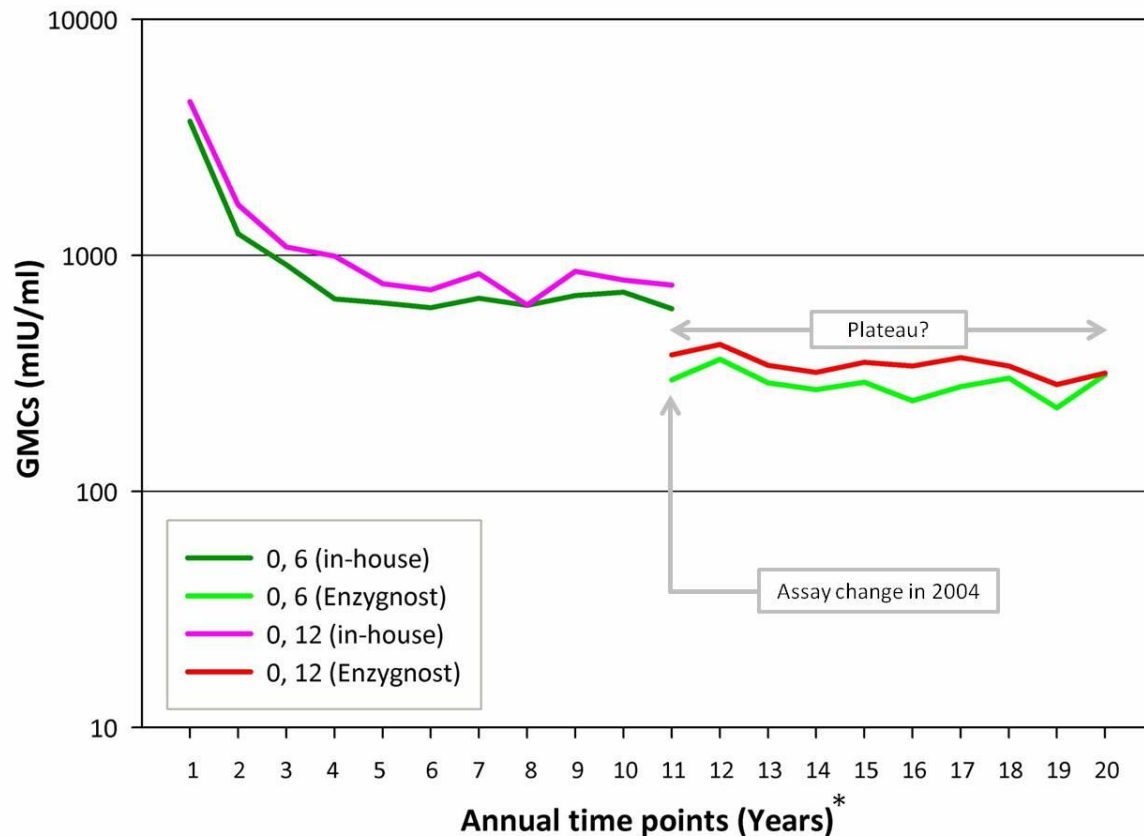
Attenuated HAV strain	Name	Adjuvant	HAV Antigen Dose/injection		Manufacturers
			Paediatric	Adult	
H2	Freeze-dried live HAV vaccine	None	0.5 ml (6.5log CCID <sub>50</sub> )	1.0 ml (6.5log CCID <sub>50</sub> )	Zhejiang Pukang Biotech company
LA-1	HAVAC Freeze-dried live HAV vaccine	None		1.0 ml (6.5log CCID <sub>50</sub> )	Changchun Institute of Biologic Products

\*\* Modified from <sup>160</sup>

# Properties of hepatitis A vaccines

- Highly immunogenic
- Flexible injection schedule
- Excellent safety record
- Long-lasting immunity
- Booster doses not required for immune-competent subjects who received two doses

# Post vaccination anti-HAV antibody levels 20 Years



- Anti-HAV antibody GMCs peaked 1 month post-dose 2 in both studies
- GMCs declined sharply during first year after primary vaccination
- Thereafter, low rate of decay in antibody levels
  - ?Plateau reached?
- 20 years post-primary vaccination anti-HAV GMCs persist at 317 mIU/ml and 312 mIU/ml in the seropositive subjects in studies HAV-112 (0, 12) and HAV-123 (0, 6), respectively

Figure provided by Pierre van Damm, Univ of Antwerp, Belgium

# Combination Vaccines

- Hepatitis A and B
  - TWINRIX®
- Hepatitis A and typhoid
  - Viatim®
  - Vivaxim®
  - Hepatryx®

# Hepatitis A Vaccines

## Control and Prevention Strategies

- Immunization of defined risk groups
- Regional mass vaccination of pediatric sub-populations at risk
- Universal vaccination of toddlers
- Single-dose immunization
- Post-exposure prophylaxis and intervention in outbreaks

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# Hepatitis A Vaccines

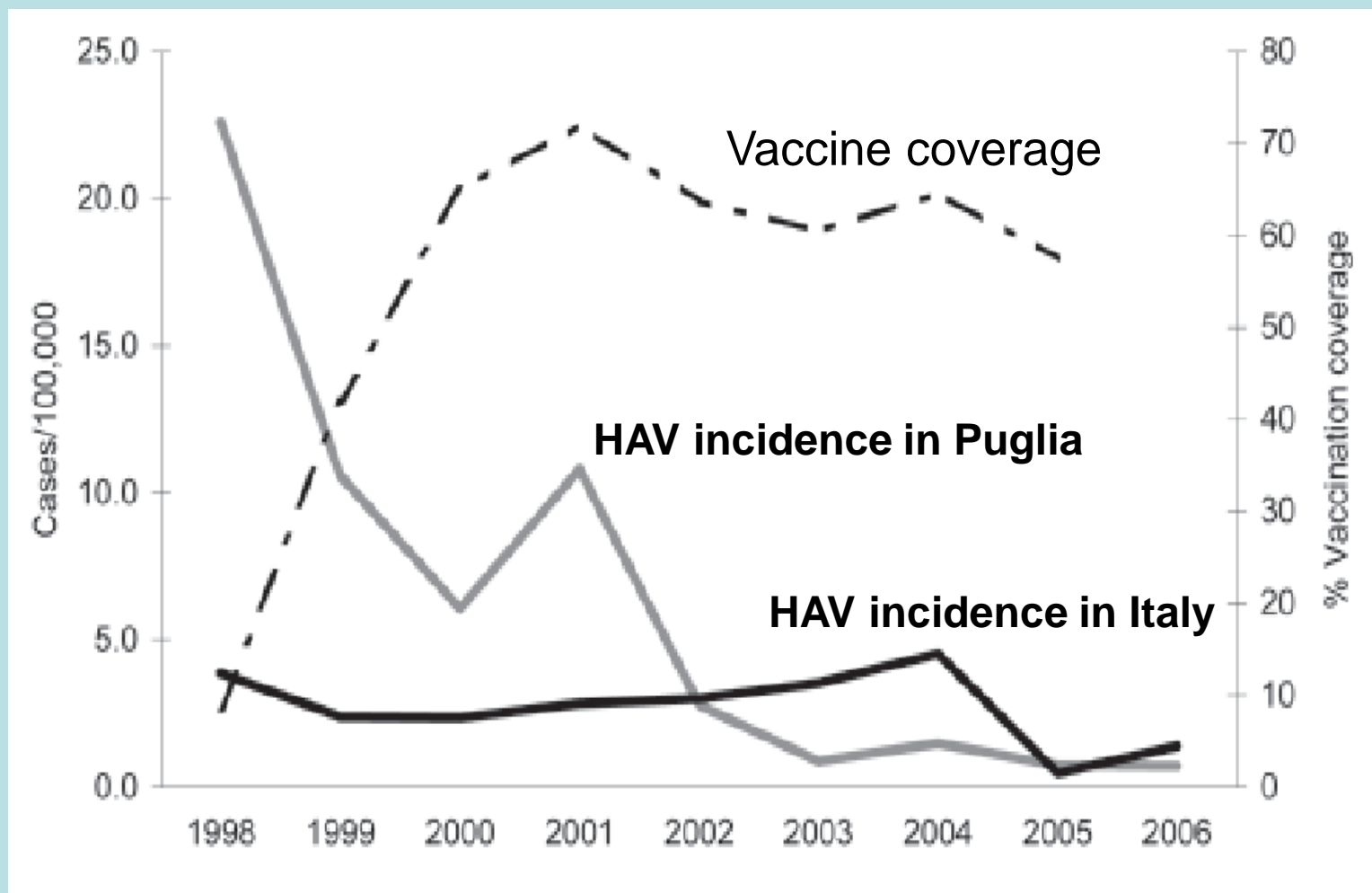
## Control and Prevention Strategies

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# Selected regional mass vaccination programs of pediatric subpopulations at risk

- US
  - Alaska
  - American Indians
  - Butte county
- Puglia, Italy
- Catalonia, Spain
- North Queensland, Australia
- Minsk, Belarus
- Shengsi county and Jiaojiang city, Zhejiang province, China (attenuated live vaccine)

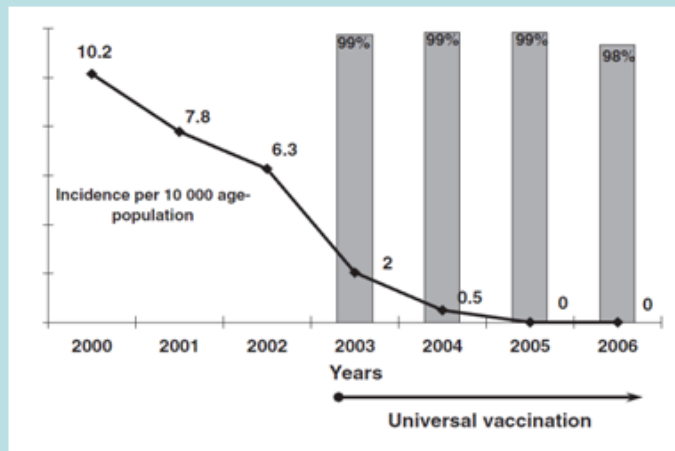
# Vaccination coverage and incidence of hepatitis A in Puglia region and Italy, 1998-2006\*



# Belarus 2003: Childhood Hepatitis A Vaccination Program in Minsk

A

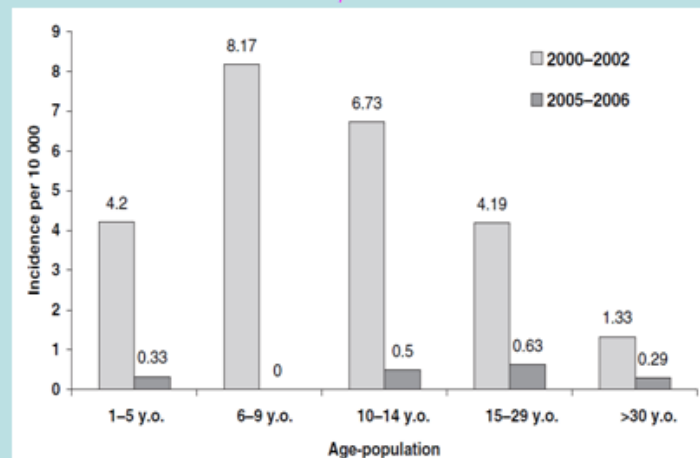
Hepatitis A Incidence in 6 Year Old Children and Vaccine Coverage, Minsk, Belarus, 2000-2006



Fisenka EG et al, J Viral Hepatitis 2008; 15 suppl 2:57

B

Hepatitis A Incidence Rate by Age Group Before and After Vaccination Program Implementation, Minsk, Belarus, 2000-2006

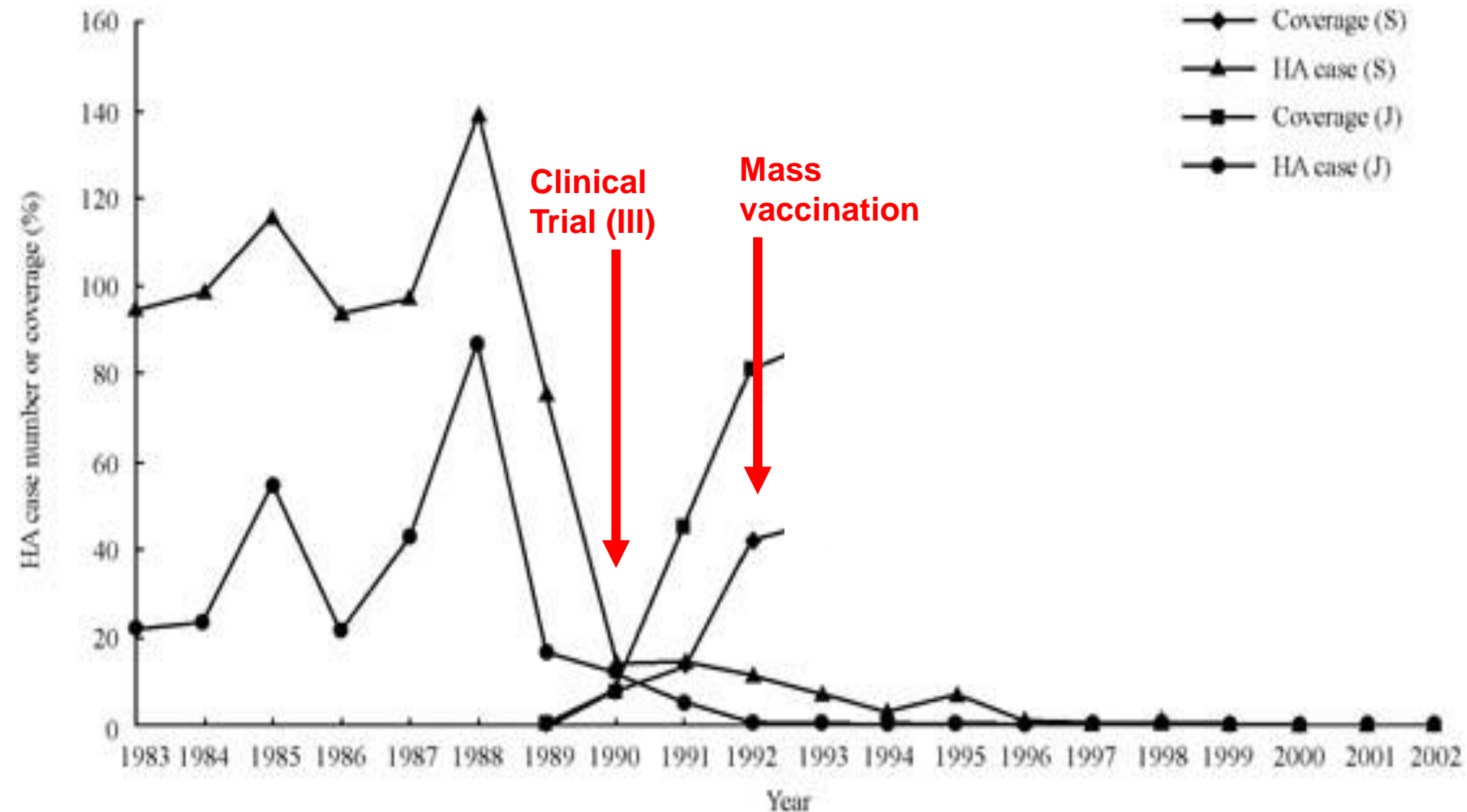


Fisenka EG et al, J Viral Hepatitis 2008; 15 suppl 2:57

# China: Hepatitis A Vaccination of Children

- Shengsi County and Jiaojiang City, Zhejiang Province, China
- Begun as demonstration project in 1992
- Initial vaccination of children ages 1-15 years
- Subsequent ongoing vaccination of each new cohort
- Single dose live attenuated vaccine (ZhePu)
- Estimated coverage 85%-91%

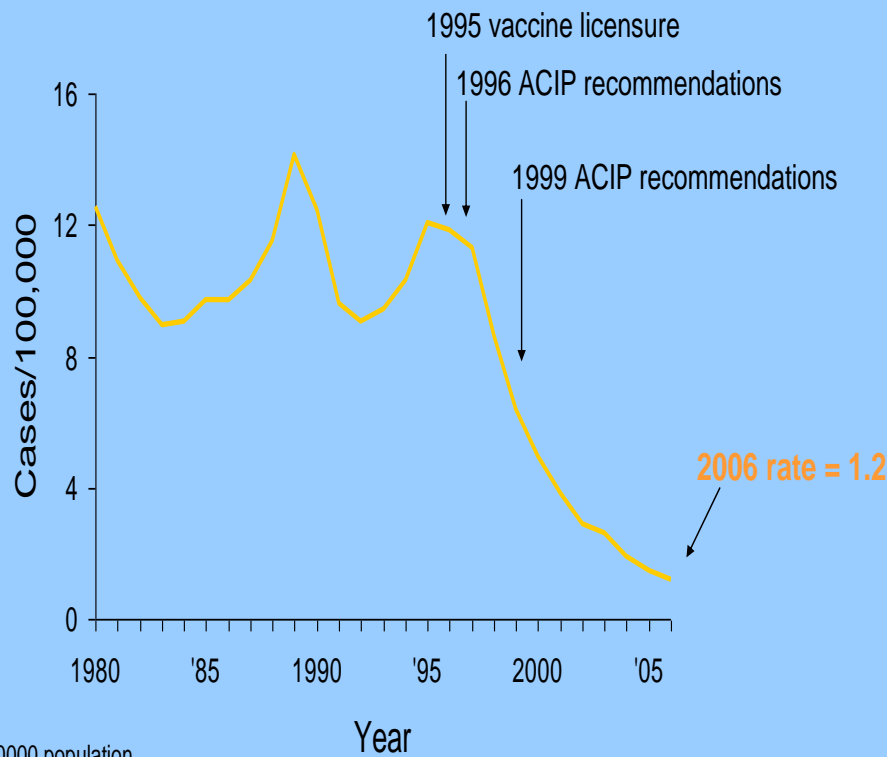
# Reported Hepatitis A Cases among Children < 16 years and Hepatitis A Vaccine Coverage, Shengsi County and Jiaojiang City, Zhejiang Province, China (1983 to 2002)



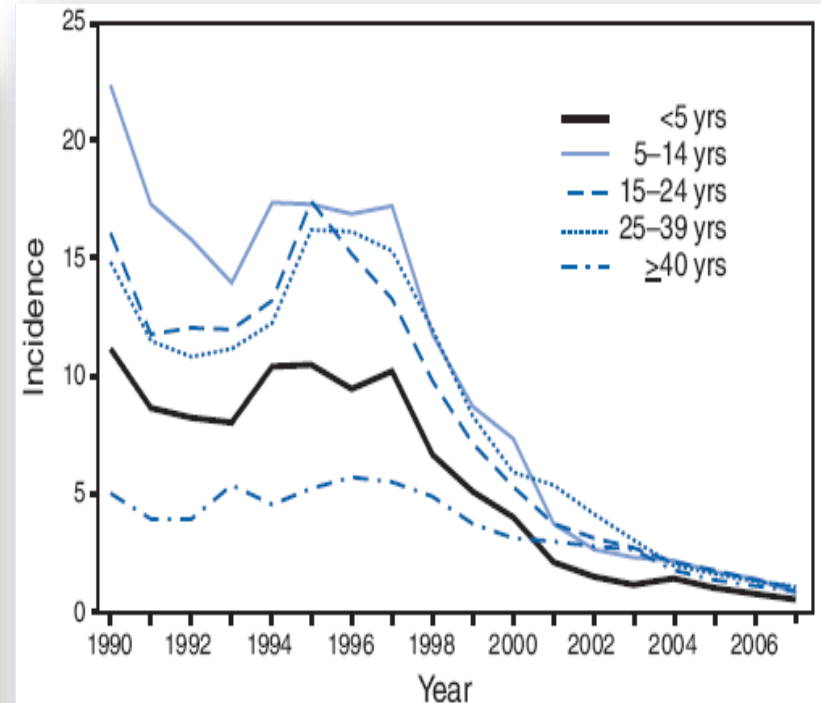
Source: Zhuang FC, et al. Chinese Medical Journal 2005;118:1851-6.

# Impact of incremental vaccination strategy against hepatitis A in the US

Overall Hepatitis A Incidence\*  
(USA, 1980-2006)



Incidence\* of acute hepatitis A, by age group and year following the gradual introduction of mass vaccination in children in the USA, 1990-2007



- One dose vaccine coverage rose from 17% to 47% between 2006-2009 in 12-23m old toddlers (8 sentinel sites).
- Full 2 dose vaccine coverage rose from 1% to 15% (MMWR 29th July, 2010)

# Impact on Health Care Utilization, U.S. 1996-2004

## Medstat MarketScan Database

Comparing baseline (1996-97) to 2004,  
statistically significant declines:

- Hospitalizations – 69%
- Ambulatory visits – 42%
- Adjusted to US population, medical expenditures for hospitalizations and ambulatory visits declined:
  - 68% reduction
  - \$29.1 million (baseline) to \$9.3 million (2004)



# Updated ACIP Recommendations for Post Exposure Prophylaxis against HAV (abbreviated)

- ➡ **For healthy persons age  $\geq$  12 months to 40 years, hepatitis A vaccine is preferred to IG.**
- ➡ **For persons > 40 years, IG is preferred.  
(Vaccine can be used if IG cannot be obtained)**
- ➡ **For children age < 12 months, immunocompromised persons, persons with chronic liver disease, and persons for whom vaccine is contraindicated, IG should be used.**

# Hepatitis A Vaccines

## Control and Prevention Strategies

- Immunization of defined risk groups
- Regional mass vaccination of pediatric subpopulations at risk
- Universal vaccination of toddlers
- Single-dose immunization
- Post-exposure prophylaxis and intervention in outbreaks

# Factors affecting vaccination strategy

- Disease burden and level of endemicity
- Socio-economic development and sanitation
- Risk of outbreaks
- Vaccine costs and cost-effectiveness
- Acceptance by the population

## Highlights of HAV Epidemiology in Israel

### Background

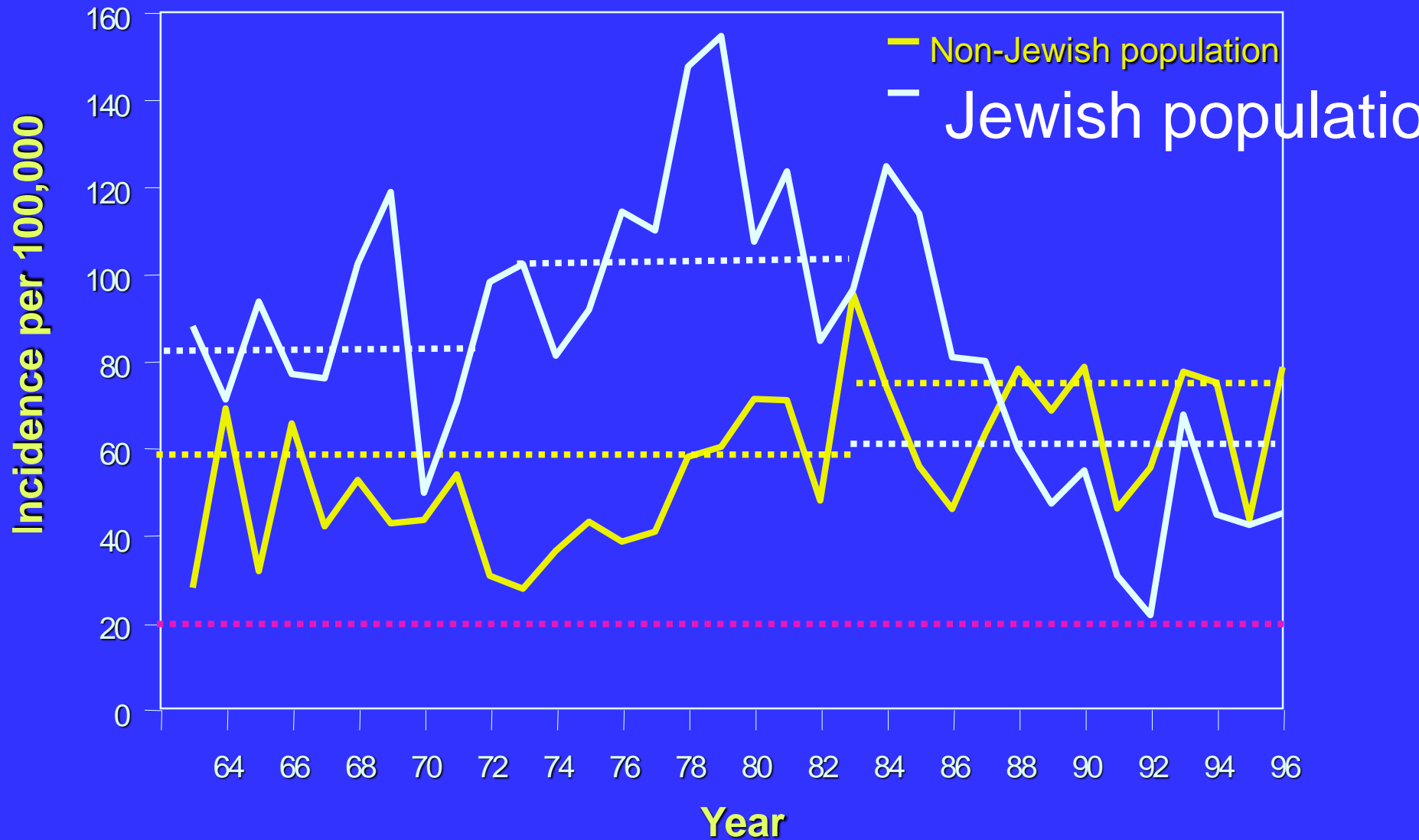
- Heterogeneous population (contact between high and low socioeconomic risk groups)
- Highest attack rate in children 5-9 years old
- Maternal anti-HAV IgG is usually cleared in babies by the age of 18 months
- Hepatitis A is rarely observed < age of 18m
- Toddlers seem to be the main vehicle for HAV transmission (pilot study results)

## Jews vs Non-Jews in Israel - Relevance to HAV Epidemiology

The non-Jewish population in Israel as compared to the Jewish population::

- Lower socioeconomic status
- Higher birthrate (37.6 vs 18.5 per 1,000)
- More crowded living conditions (2.99 vs 2.18 per household)
- Lower hygienic infra-structure

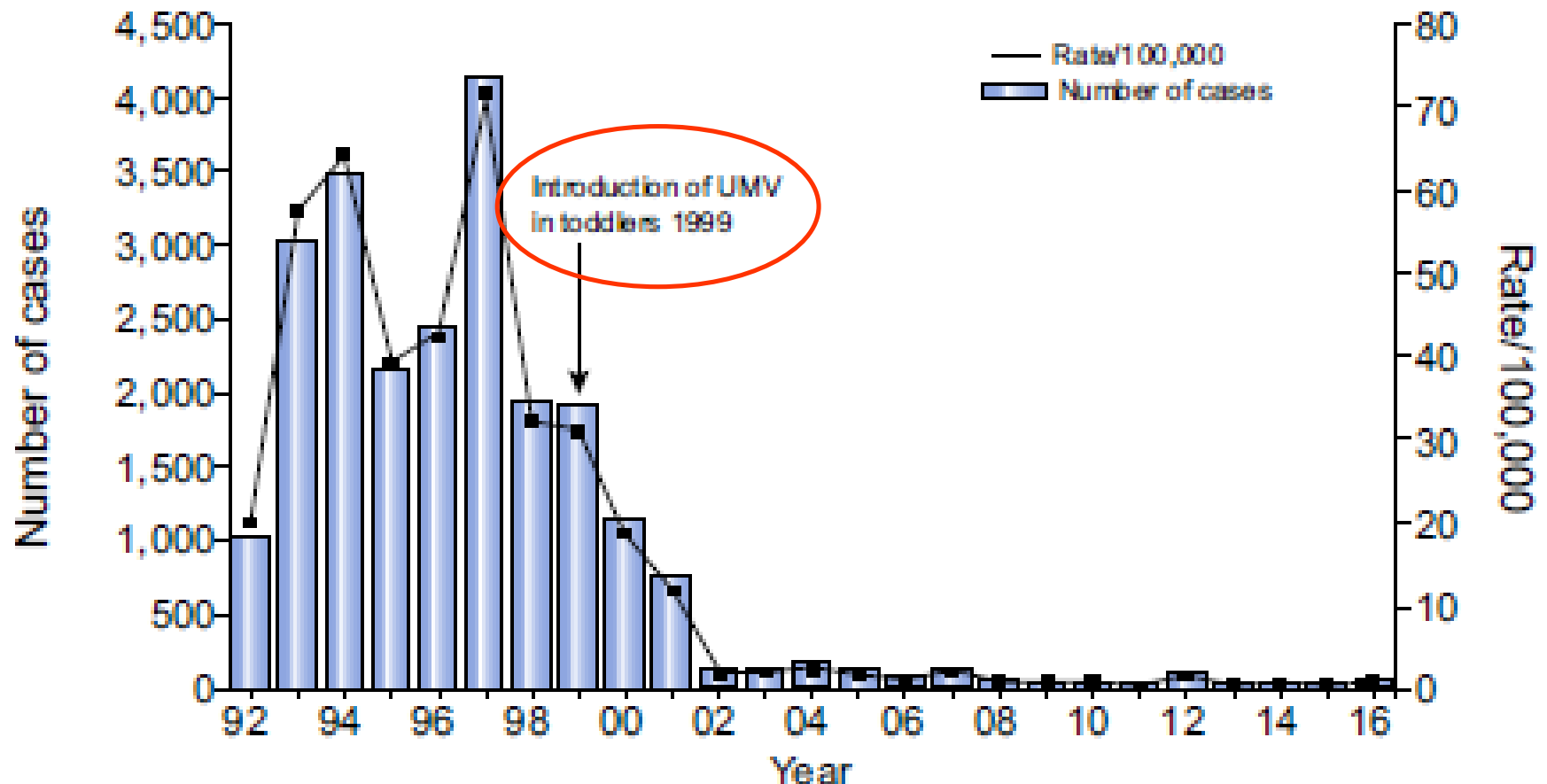
# Incidence of Viral Hepatitis in Israel 1963-1996 by Population



# Israel: Childhood Hepatitis A Vaccination Program

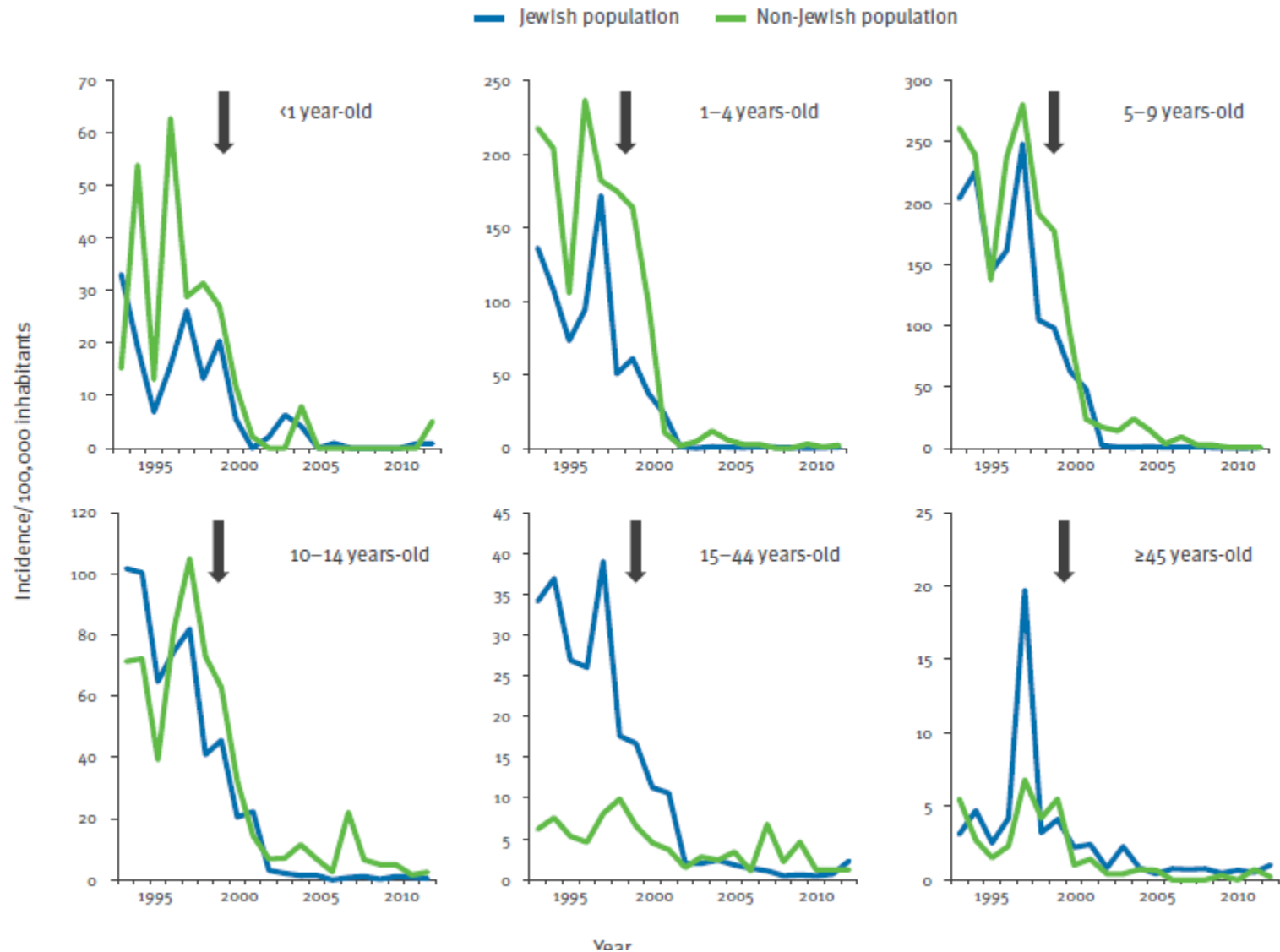
- July 1999
- Vaccination of all 18 month old children
- Vaccine provided free of charge, as part of regular immunization program
- Estimated first dose coverage in vaccinated cohorts – 90%; second dose – 85%

# Incidence of Hepatitis in Israel 1996-2016



Lemon SM, Ott JJ, Van Damme P, Shouval D. J Hepatol. 2017

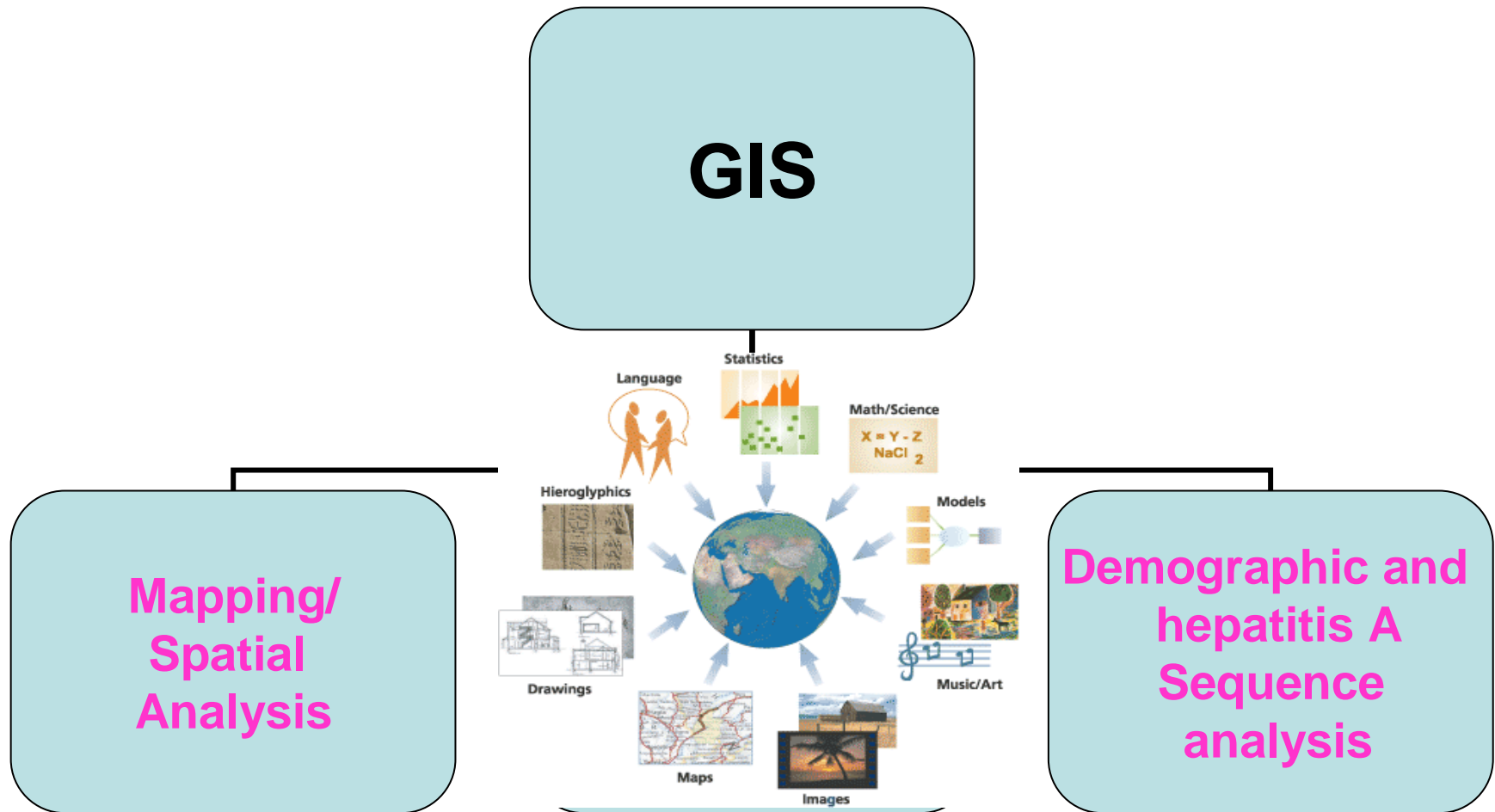
# Annual incidence rates of hepatitis A/100,000 population, by specific age and ethnicity, 1993-2012





# Geographic Information System (GIS)

A system of **hardware** and **software** linking **mapped objects** to **collected** information( i.e. epidemiologic data)

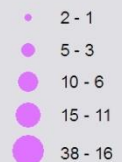


# HAV isolates in Jerusalem

## Cluster 1a



## Cluster 1b



### Population type



0 2,500 5,000  
Meters

Produced: The GIS Center, HUJI  
Israel 2007

# Number of Acute Hepatitis A Cases in the Jerusalem District\*

**1999      671**

**2000      654**

**2001      420**

**2002      46**

**2003      67**

**2004      50**

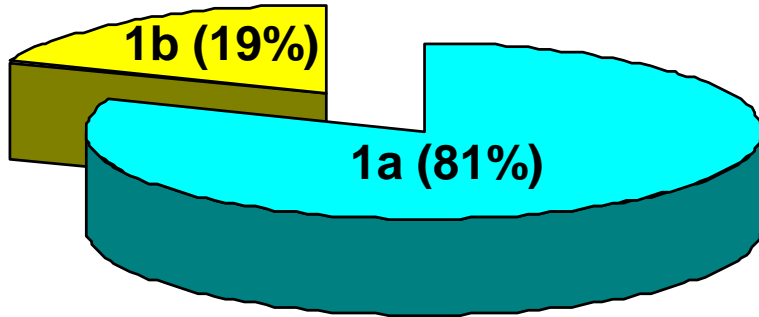
**total      1908**

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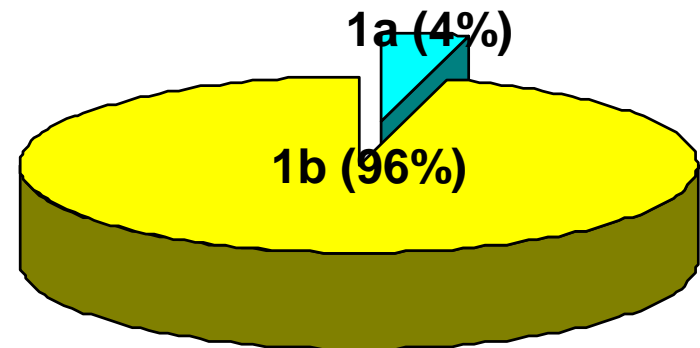
\*Through active surveillance in a population of  
~900,000

# HAV Genotype Distribution in Sub-populations - Jerusalem

**Jewish population (N=466)**

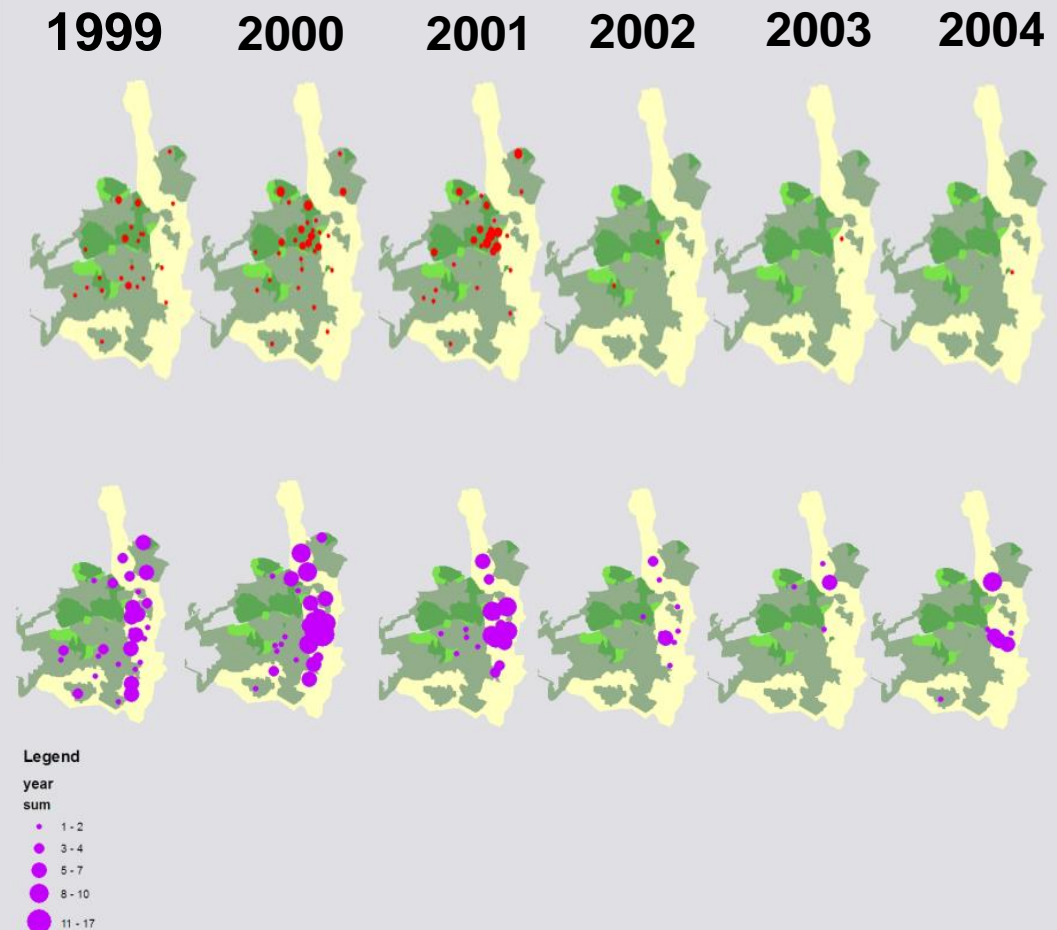
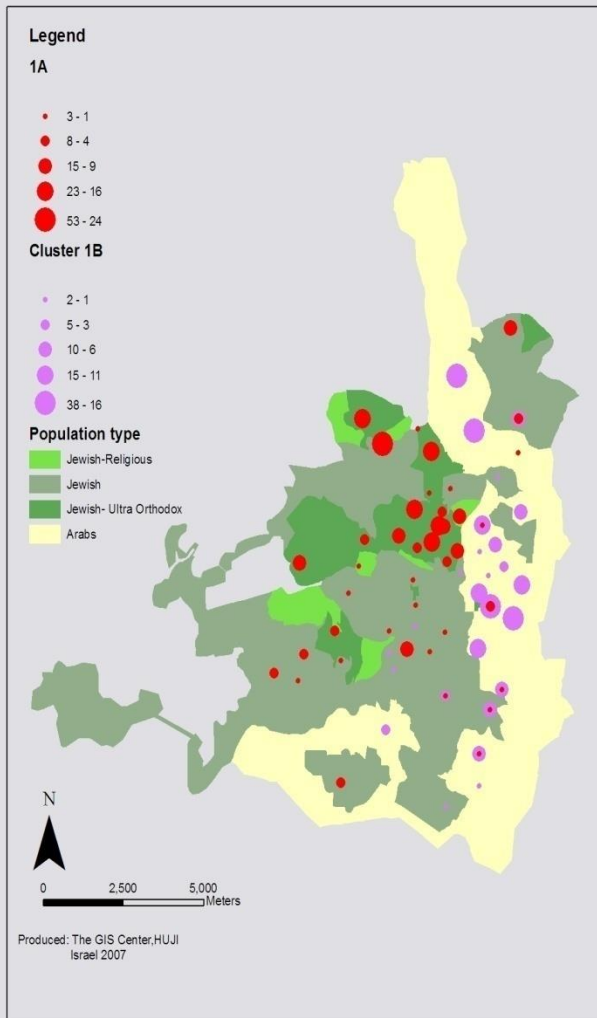


**Non-Jewish population (N=243)**

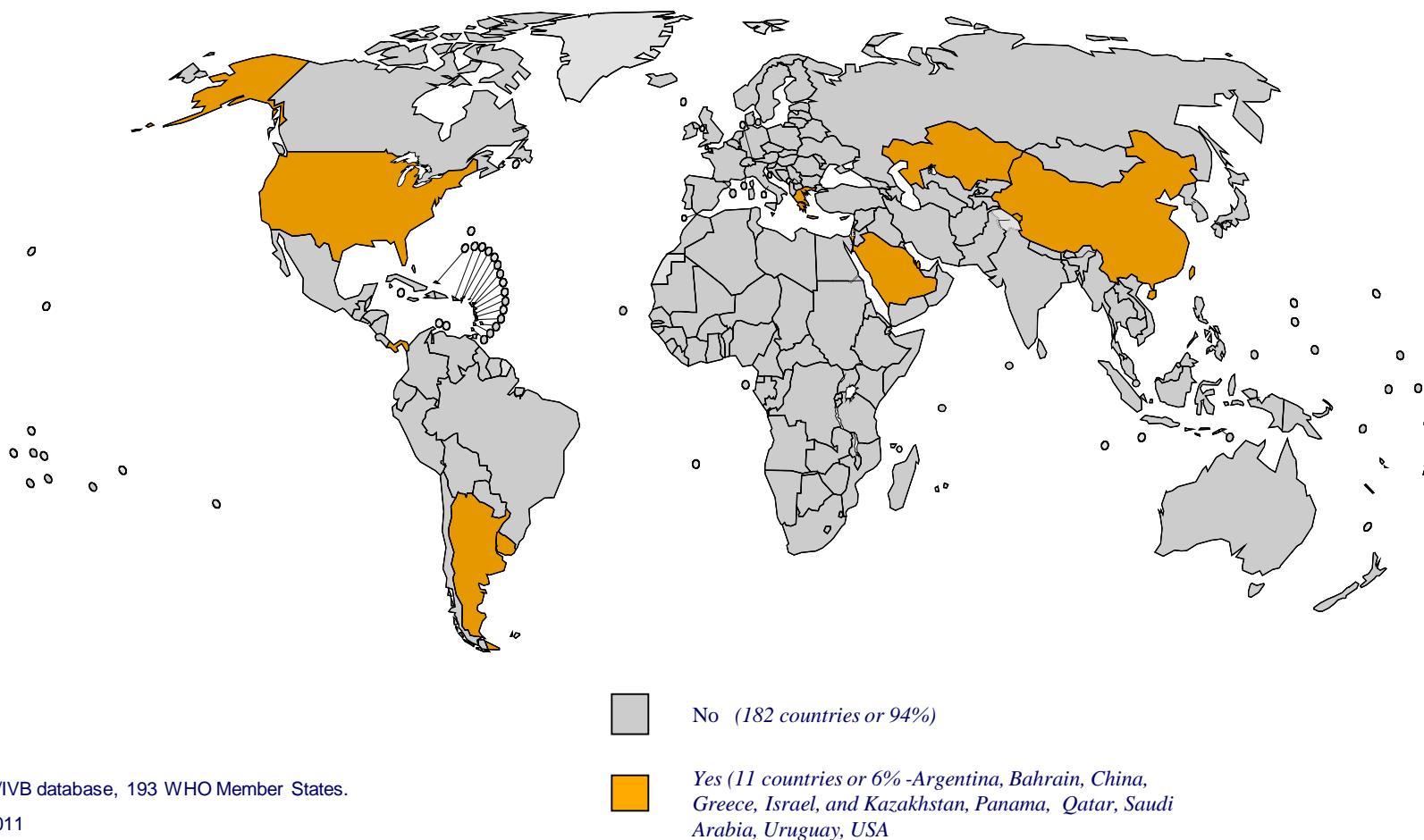


# - Control of HAV infection in Jerusalem

## - Follow-up on GIS



# Countries Using HepA Vaccine in National Immunization Schedule, 2010



Source: WHO/IVB database, 193 WHO Member States.

27 October 2011

The boundaries and names shown and the designations used on this map do not imply the expression of any opinion whatsoever on the part of the World Health Organization concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted lines on maps represent approximate border lines for which there may not yet be full agreement. © WHO 2011. All rights reserved

# Impact of universal mass vaccination with monovalent inactivated hepatitis A vaccines - A systematic review

The WHO recommends integration of universal mass vaccination (UMV) against HAV in national immunization schedules for children aged  $\geq 1$  year, if justified on the basis of acute HAV incidence, declining endemicity from high to intermediate and cost-effectiveness. This recommendation has been implemented in several countries.

## **-Review of 27 studies (Argentina, Belgium, China, Greece, Israel, Panama, the United States and Uruguay).**

- All except one study showed a marked decline in the incidence of hepatitis A post introduction of UMV.
- The incidence in non-vaccinated age groups decreased as well, suggesting herd immunity but also rising susceptibility.
- Long-term anti-HAV antibody persistence was documented up to 17 y after a 2-dose primary vaccination.

**Conclusion:** introduction of UMV in countries with intermediate endemicity for HAV infection led to a considerable decrease in the incidence of hepatitis A in vaccinated and in non-vaccinated age groups alike

# Hepatitis A Vaccines

## Control and Prevention Strategies

- Immunization of defined risk groups
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# Fulminant hepatitis A in children

Number of reports is rising?

- Turkey 4 cases (6/04-11/06)
- UK 9 cases (1991-2000))
- Argentina 128 cases (5/82-9/02)  
41 cases (9/03-1/06)
- Brazil 13 cases (1998-2007)
- Korea 35 cases (2003-2008)

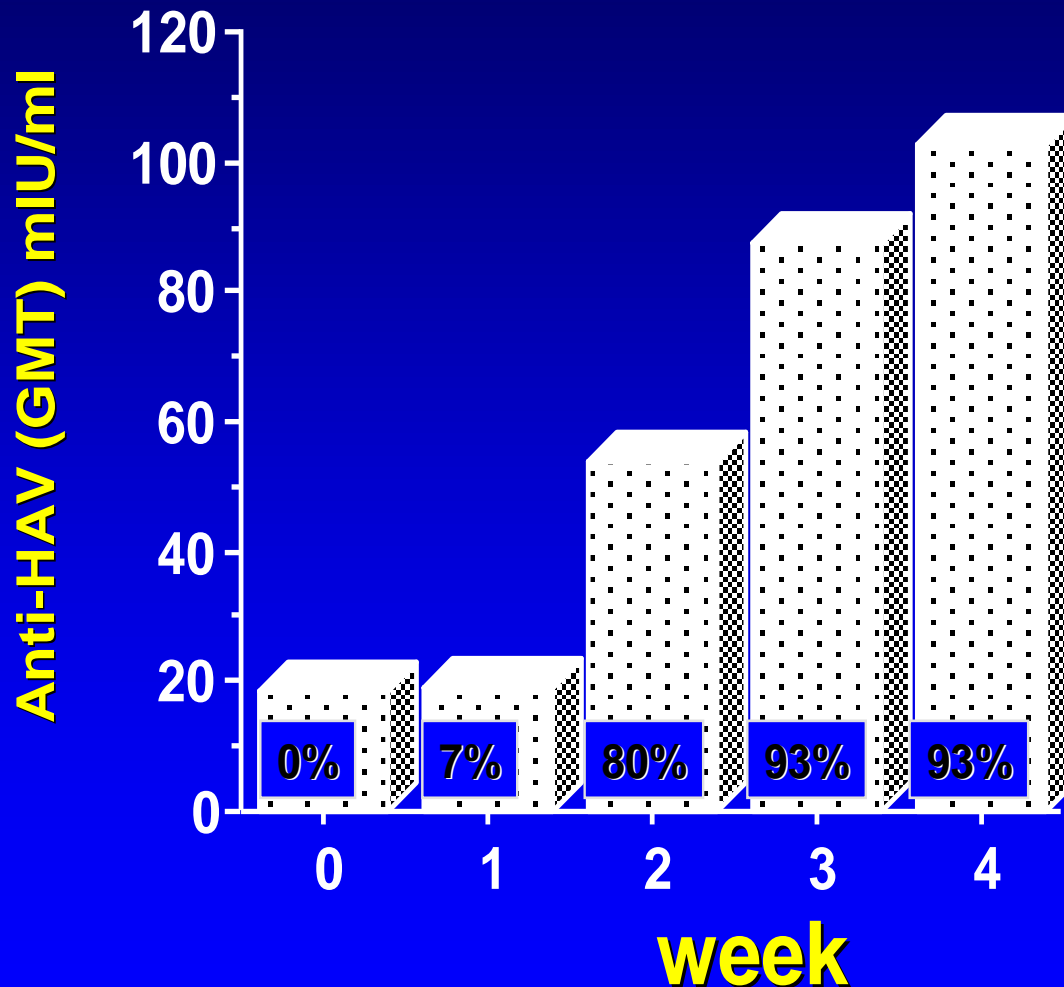
***Reports are retrospective and released by individual centers***

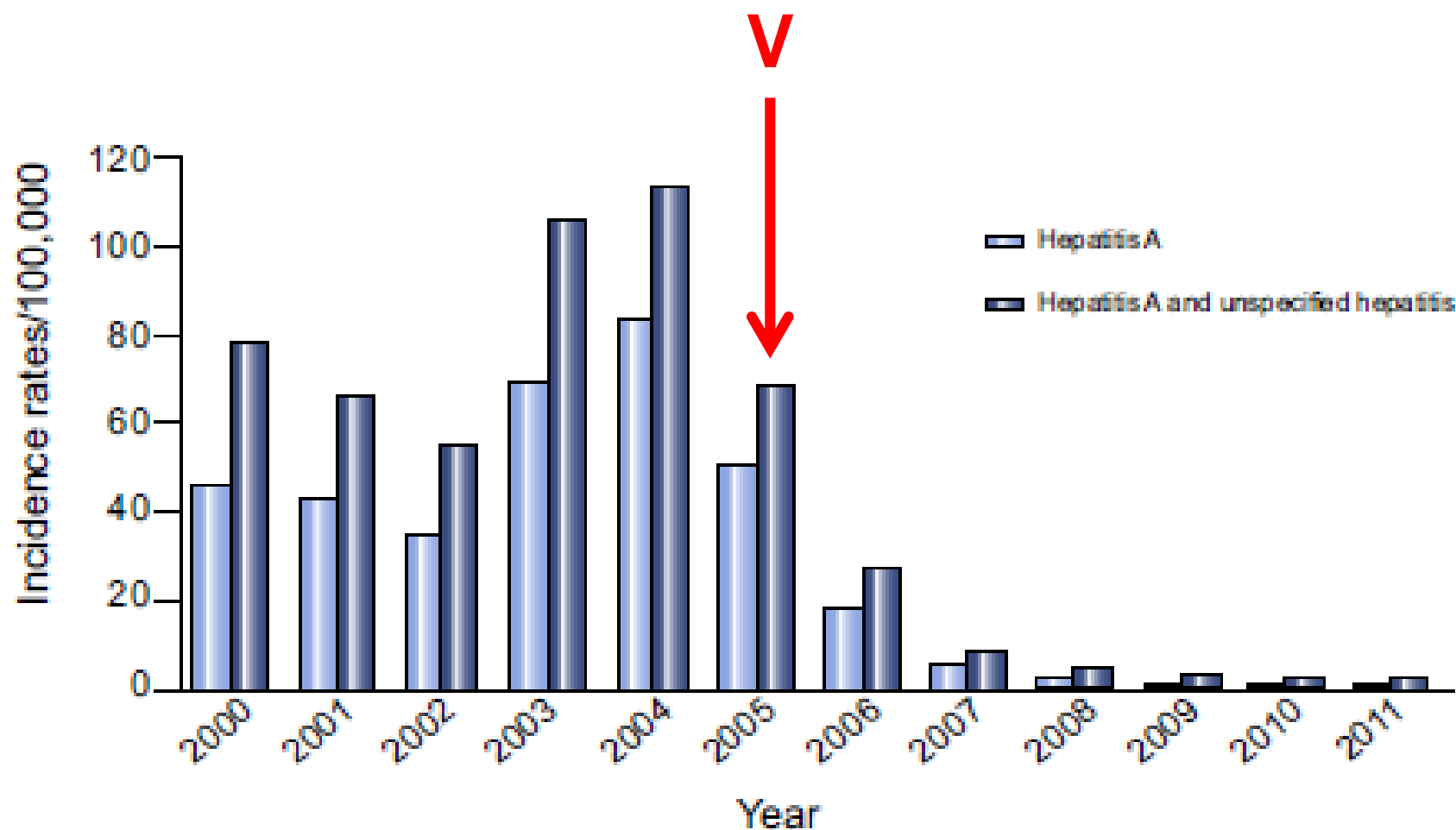
*J Viral Hepatitis 2008;15:S66; J Ped Gastroenterol & Nutr 2005;40:575; Pediat Crit Care Med 2002;3:227; Liver Int 2007;Arch Dis Child 2008;93:48*

# Argentina: Childhood Hepatitis A Vaccination Program

- Universal single-dose hepatitis A immunization program
- June 2005
- Children aged 12 months
- Most vaccines provided free of charge
- Vaccine coverage 95% in 2006
- 80% decrease in incidence from 70.5-173.8/100,000 to 10.2/100,000

# Rapid Seroconversion Following a Single Dose of an HAV Vaccine





**Fig. 8. Impact of the single-dose immunisation strategy against hepatitis A in Argentina.** (Reproduced with permission from<sup>177</sup>).

# Hepatitis A Vaccines

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# Hepatitis A vaccine versus immune globulin for post-exposure prophylaxis

- 1090 household and day-care contacts , 2-40y old of index cases randomized to receive an hAV vaccine or IG
- Transmission of HAV confirmed by anti-HAV IgM occurred in 4.4% of vaccine and 3.3% IG recipients (RR 1.35;95% CI:0.70-2.67)

# Key Concepts

## Hepatitis A is a vaccine preventable disease

- ✓ Hepatitis A virus infection is a self limited disease but it still causes significant morbidity in young and older adults, associated with temporary disability and cost
- ✓ Large populations of adolescents and young adults in countries with intermediate endemicity (and in transition”) who escaped HAV infection in their early childhood are at risk for contracting clinical hepatitis A due to the current shift in susceptibility. This trend may lead to potential outbreaks
- ✓ Immunization of defined risk groups has a limited impact on overall burden of infection
- ✓ Universal immunization against hepatitis A to babies is highly effective in controlling transmission to children and provides herd immunity to unvaccinated adults
- ✓ Booster dose(s) are not required after successful immunization
- ✓ Post exposure prophylaxis using an hepatitis A vaccine within 14 days of exposure, is effective with an important advantage of providing much longer protection against hepatitis A as compared to immune globulin

# Two outbreaks of HAV

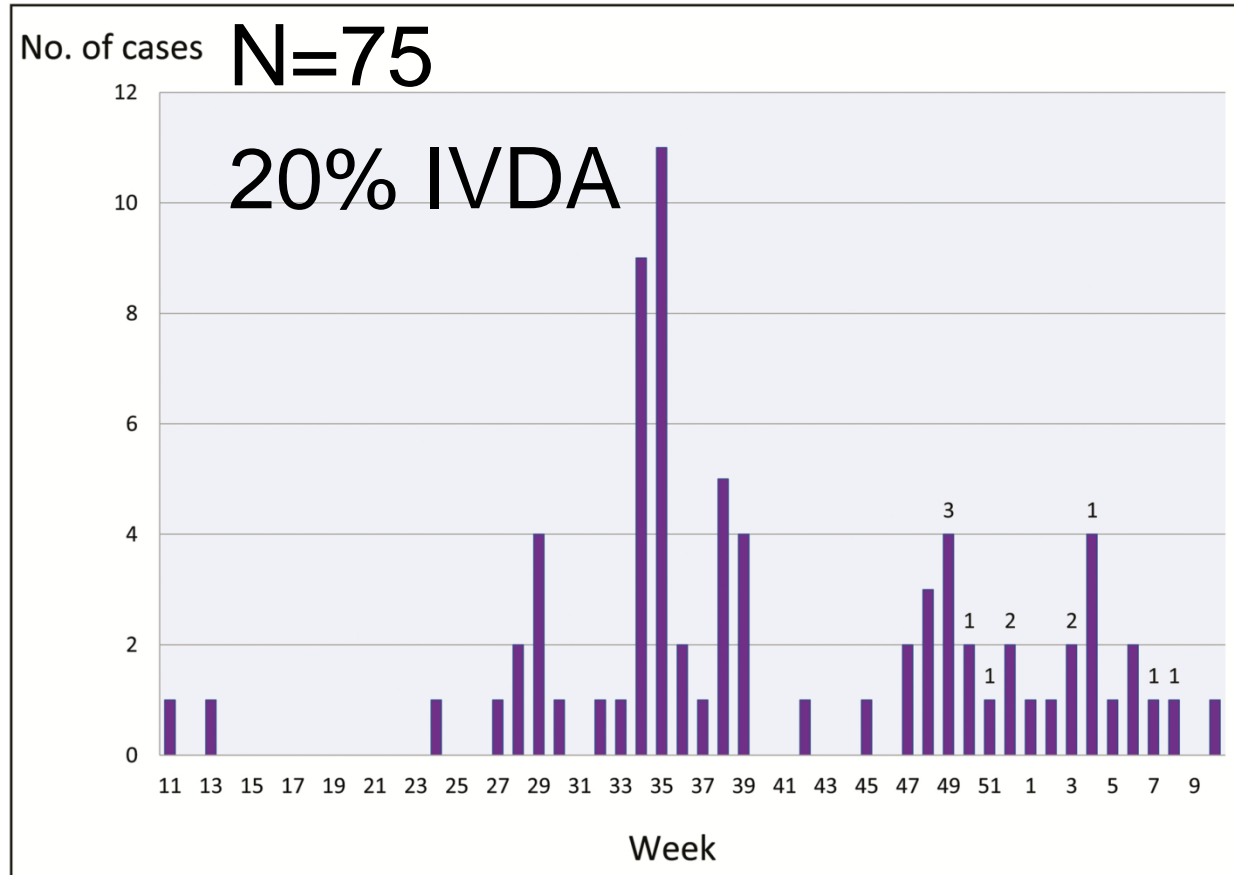
- IVDA
- MSM

Evidence for Hepatitis A Virus Endemic Circulation in Israel Despite Universal Toddler Vaccination Since 1999 and Low Clinical Incidence in All Age Groups

Yosef Manor,<sup>1</sup> Matthew Lewis,<sup>2</sup> Daniela Ram,<sup>1</sup> Nili Daudi,<sup>3</sup> Orna Mor,<sup>1</sup> Michal Savion,<sup>2</sup> Zipi Kra-Oz,<sup>4</sup> Yonat Shemer Avni,<sup>5</sup> Rivka Sheffer,<sup>2</sup> Daniel Shouval,<sup>3</sup> and Ella Mendelson<sup>1,6</sup>



# Outbreak of HAV in the Tel Aviv District 2012-2013



# HAV clinical cases, 2017

- Number of reported cases: 81
- Median age: 34y (range 3-56y)
- Male/Female: 69/24 (85.2% M)

# Sewage derived environmental surveillance

- A useful tool for population-based monitoring of microbial and viral activities.
- In Israel, a national program for surveillance of the poliovirus in sewage has been ongoing since 1988. It led to the identification of wild poliovirus in 2013, which activated a major emergency response by the Public Health Services
- Similar surveillance is currently employed for HAV

# Sampling of Urban Sewage



Shafdan STF

500 ml sample(filter& centrifuge) to 15m



Extract NA from  
1ml



Real-Time PCR  
(TQM)



HAV negative



**HAV positive**



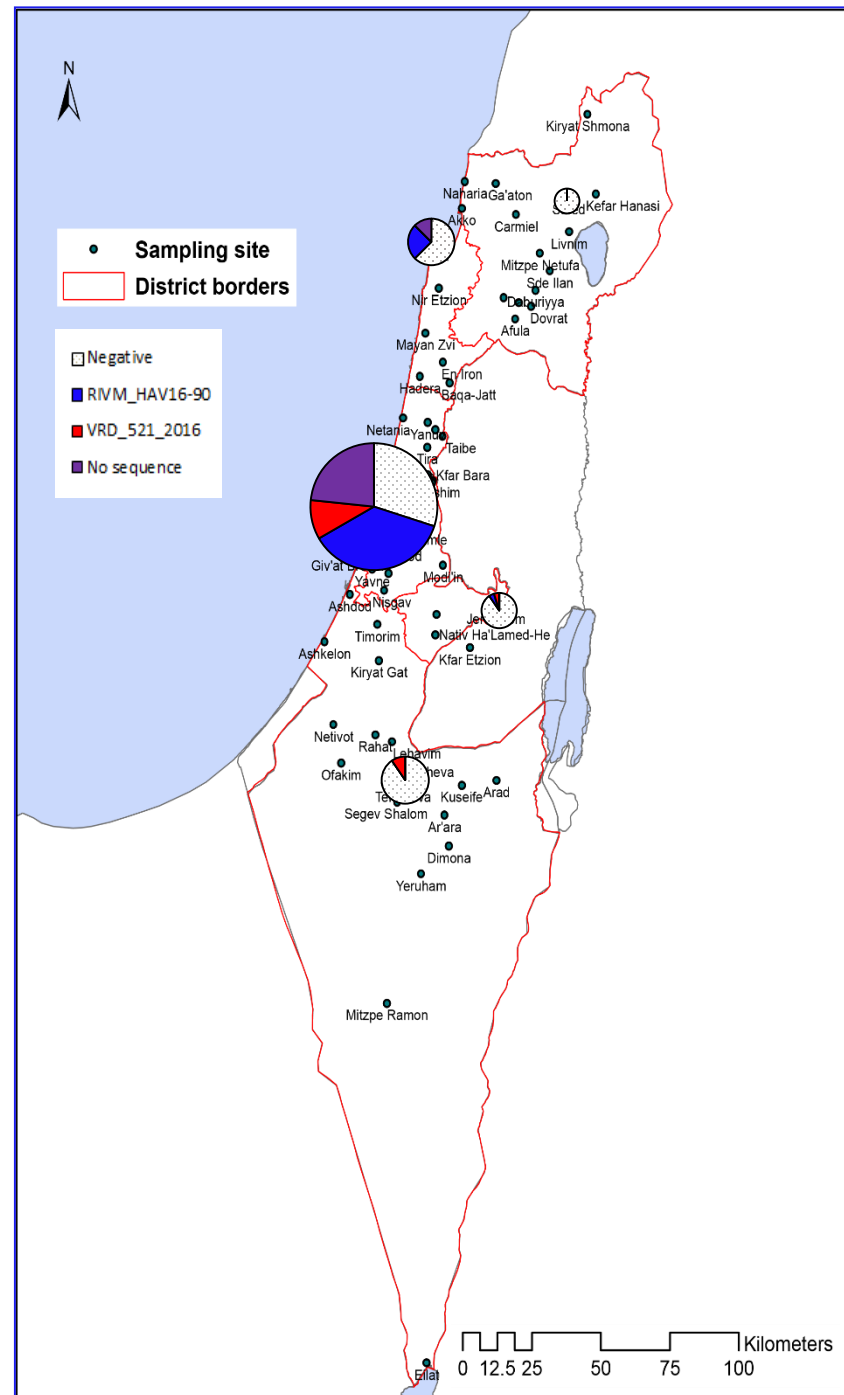
sequence



# Environmental surveillance

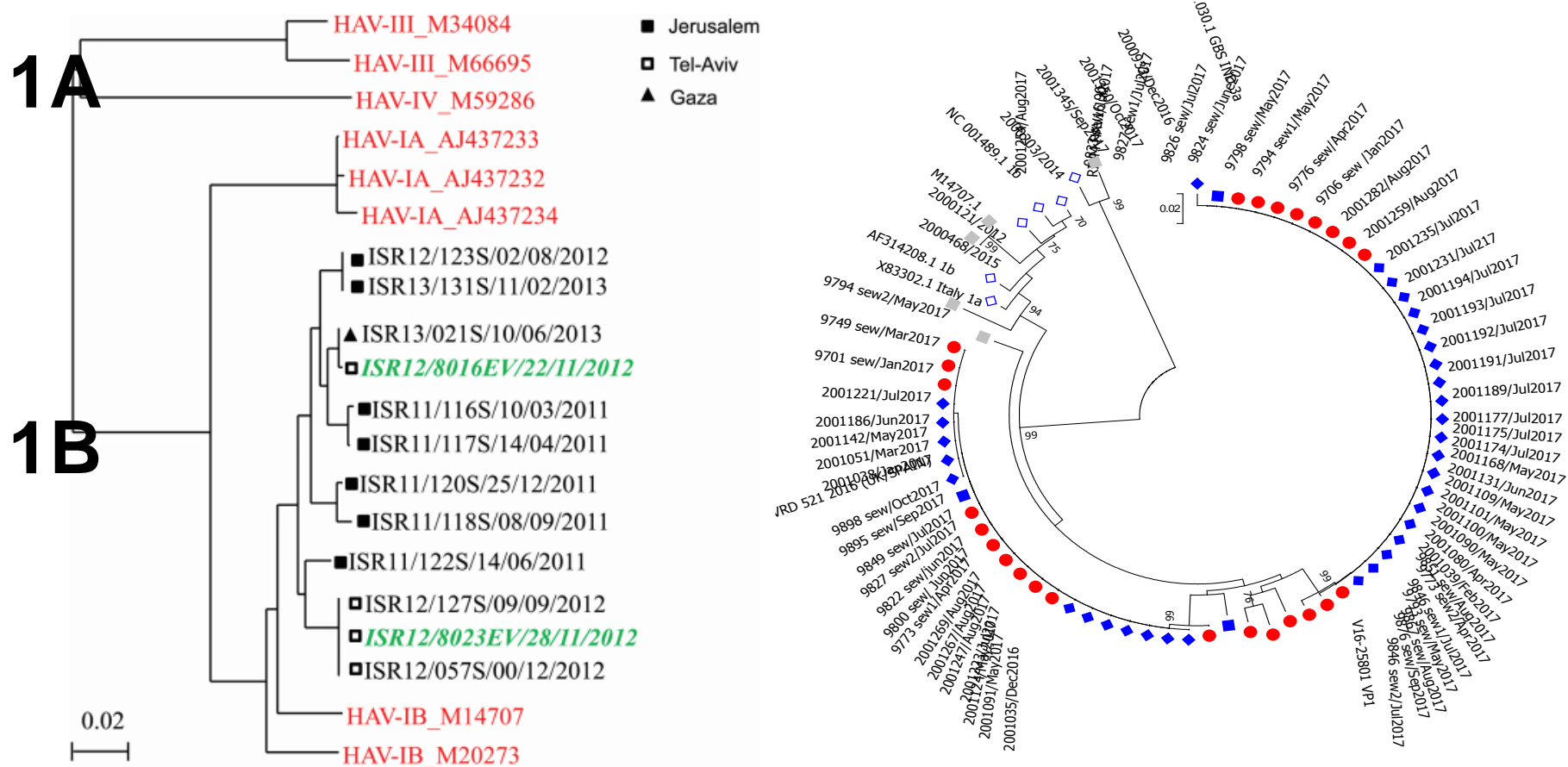
147 sewage samples were collected monthly from 14 facilities around the country

31% (45/147) of sewage samples were HAV positive, with a high prevalence (63%, 26/41) in facilities in the Tel-Aviv area.



Phylogenetic tree showing the Israeli (black and green ) and the reference (red) strains belonging to genotypes IA, IB, III, and IV, as indicated in each reference strain name.

Black letters indicate serum samples, and green i indicate environmental samples.



# Summary

- The HAV 1a outbreak in MSM in Israel is m.p. imported , from European countrise. HAV 1B in IVDA is mainly derived from the Gaza strip and possibly Jordan West bank
- 
- Despite the efficient universal mass vaccination program which lead to a dramatic fall in the annual HAV incidence (from 33-70 cases to 2,5 cases/100,000), HAV can still be transmitted to susceptible/ high-risk adult population, raising the issue of catch-up vaccination.
- The role of environmental sampling in disease surveillance is demonstrated.

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<sup>1</sup>Central virology lab, MOH , Sheba Medical Center, <sup>2</sup>Central Virology Lab, Ministry of Health, <sup>3</sup>Central Virology Lab, MOH, Sheba Medical Center, Ramat-Gan, <sup>4</sup>Health district Officer, Ministry of Health, Tel-Aviv, <sup>5</sup>Epidemiology, MOH Public Health, Jerusalem, <sup>6</sup>Epidemiology, MOH Public Health, Tel-Aviv, <sup>7</sup>Liver Unit, Sheba Medical Center, Ramat-Gan, <sup>8</sup> Liver Unit, HADASSAH HEBREW UNIVERSITY HOSPITAL, LIVER UNIT, Jerusalem, Israel

Thank You



# Immunization, Vaccines and Biologicals



World Health  
Organization

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- WHO position paper on hepatitis A vaccines – June 2012 No. 28-29, 2012, 87, 261–276 <http://www.who.int/wer>