

Declaración de potenciales conflictos de intereses

Impacto de la vacunación frente a rotavirus

Prof. Dr. Federico Martinón-Torres

Relativas a esta presentación existen las siguientes relaciones que podrían ser percibidas como potenciales conflictos de intereses:

Ha recibido honorarios por conferencias de Abbot, GSK, Sanofi Pasteur MSD, Wyeth/Pfizer, y Novartis.

Ha recibido becas/ayudas de investigación de Sanofi Pasteur MSD, Wyeth/Pfizer, y Novartis.

Ha recibido honorarios por consultorías para Astra-Zeneca, Wyeth/Pfizer, Novartis, Medimmune y Sanofi Pasteur MSD.

Participa / Coordina ensayos clínicos de vacunas de Sanofi Pasteur MSD, Sanofi, Wyeth, Merck, Pfizer, Roche, Medimmune, Novartis, and GSK.



X Jornadas de Actualización en
Vacunas

Almería, 24-26 octubre de 2013

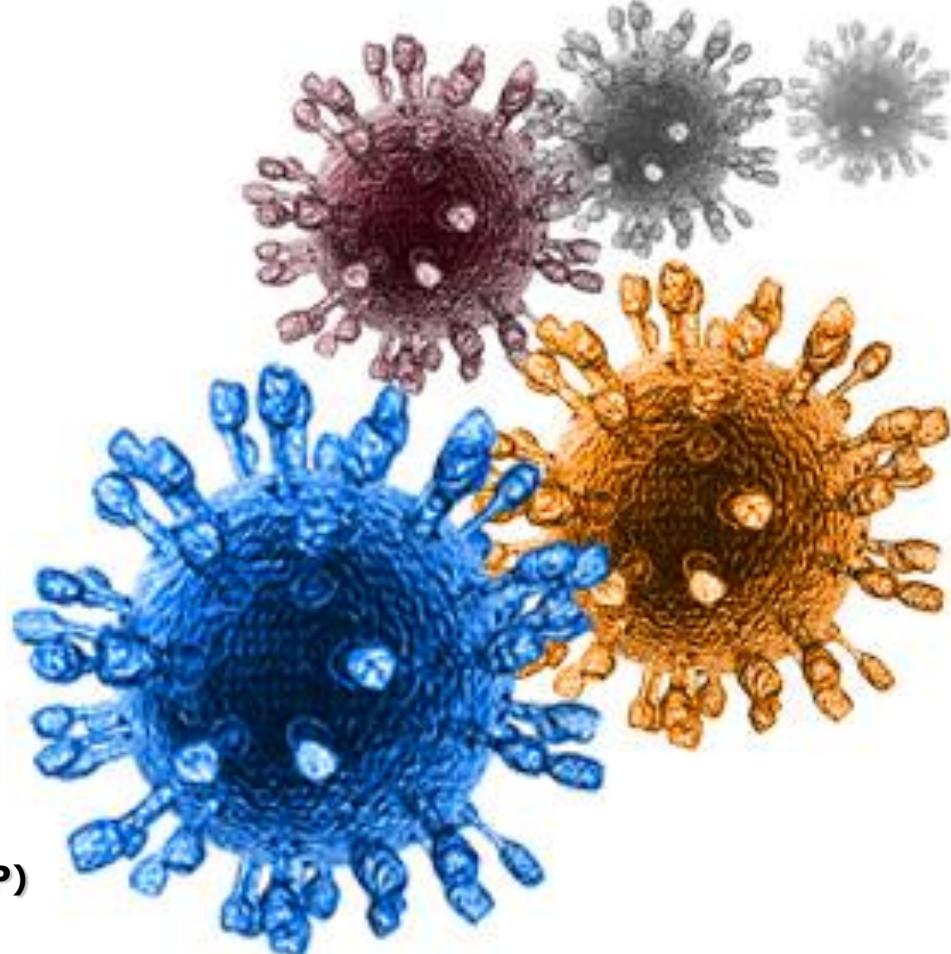
Impacto de la vacunación frente a rotavirus

Prof. Dr. F. Martinón-Torres

Pediatra, Hospital Clínico Universitario de Santiago

Grupo de Genética, Vacunas, Infecciones y Pediatría (GENVIP)

Instituto de Investigación Sanitaria de Santiago (IDIS)

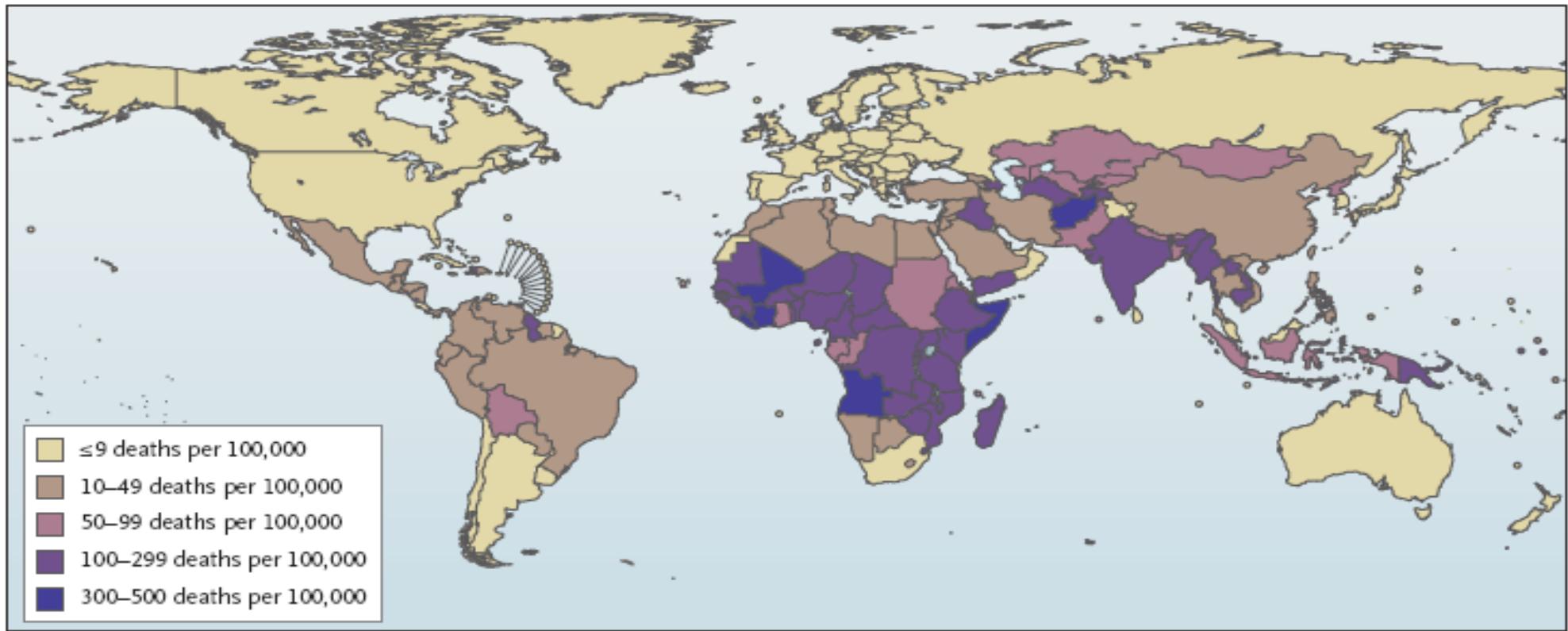


Defeating Rotavirus? The Global Recommendation for Rotavirus Vaccination

Margaret H. Danchin, M.D., Ph.D., and Julie E. Bines, M.D.



N ENGL J MED 361;20 NEJM.ORG NOVEMBER 12, 2009



Mortality Rate Associated with Rotavirus Disease.

Data are from the World Health Organization.

@fedemartinon, Almería 2013

Safety and Efficacy of a Pentavalent Human–Bovine (WC3) Reassortant Rotavirus Vaccine

Timo Vesikari, M.D., David O. Matson, M.D., Ph.D., Penelope Dennehy, M.D., Piero Van Damme, M.D., Ph.D., Mathuram Santosham, M.D., M.P.H., Zohreh Dabbaghzadeh, M.D., Michael L. Dallas, Ph.D., Jeanne M. Hayes, Ph.D., Michelle G. Goveas, M.D., L.P.H., Steve B. Eick, M.D., Mark V. Shields, M.D., Celia D.C. Christie, M.D., M.P.H., Samuli Ylitalo, M.D., Robert P. Izquierdo, Ph.D., Michele L. Coia, B.A., Matthew T. Onorato, B.S., Ben A. Adeyi, M.P.H., Gary S. Marshall, M.D., Leif Gothe fors, M.D., Dirk Campens, M.D., Aino Karvonen, M.D., James P. Watt, M.D., M.P.H., Katherine L. O'Brien, M.D., M.P.H., Mark J. DiNubile, M.D., H Fred Clark, D.V.M., Ph.D., John W. Boslego, M.D., Paul A. Offit, M.D., and Penny M. Heaton, M.D., for the Rotavirus Efficacy and Safety Trial (REST) Study Team

N Engl J Med 2006;354:11-22.

95.8% [90–98.2%]

Safety and Efficacy of an Attenuated Vaccine against Severe Rotavirus Gastroenteritis

Guillermo M. Ruiz-Palacios, M.D., Irene Pérez-Schael, M.Sc., F. Raúl Velázquez, M.D., Hector Abate, M.D., Thomas Becker, M.D., Luis Alain Costa-Gremillion, M.D., Brigitte Cheuvart, M.D., Leon Espinosa, M.D., Paul Gillard, M.D., Bruce L. Johnson, M.D., Marisol L. Llanos-Cuentas, M.D., Alexander C. Lloberas, M.D., Ricardo López, M.D., Mercedes Mayorga-Paz, M.D., Eduardo Montegut Carríao, M.D., Vest R. Richardson, M.D., Doris María Alvarez-Velarde, M.D., Luis Villera, M.D., Belén Salinas, M.D., Noris Pavía-Ruz, M.D., Jorge Salmerón, M.D., Ricardo Rüttimann, M.D., Juan Carlos Tinoco, M.D., Pilar Rubio, M.D., Ernesto Nuñez, M.D., M. Lourdes Guerrero, M.D., Juan Pablo Yarzábal, M.D., Silvia Damaso, M.Sc., Nadia Tornieporth, M.D., Xavier Sáez-Llorens, M.D., Rodrigo F. Vergara, M.D., Timo Vesikari, M.D., Alain Bouckenooghe, M.D., Ralf Clemens, M.D., Ph.D., Béatrice De Vos, M.D., and Miguel O'Ryan, M.D., for the Human Rotavirus Vaccine Study Group*

N Engl J Med 2006;354:23-33.

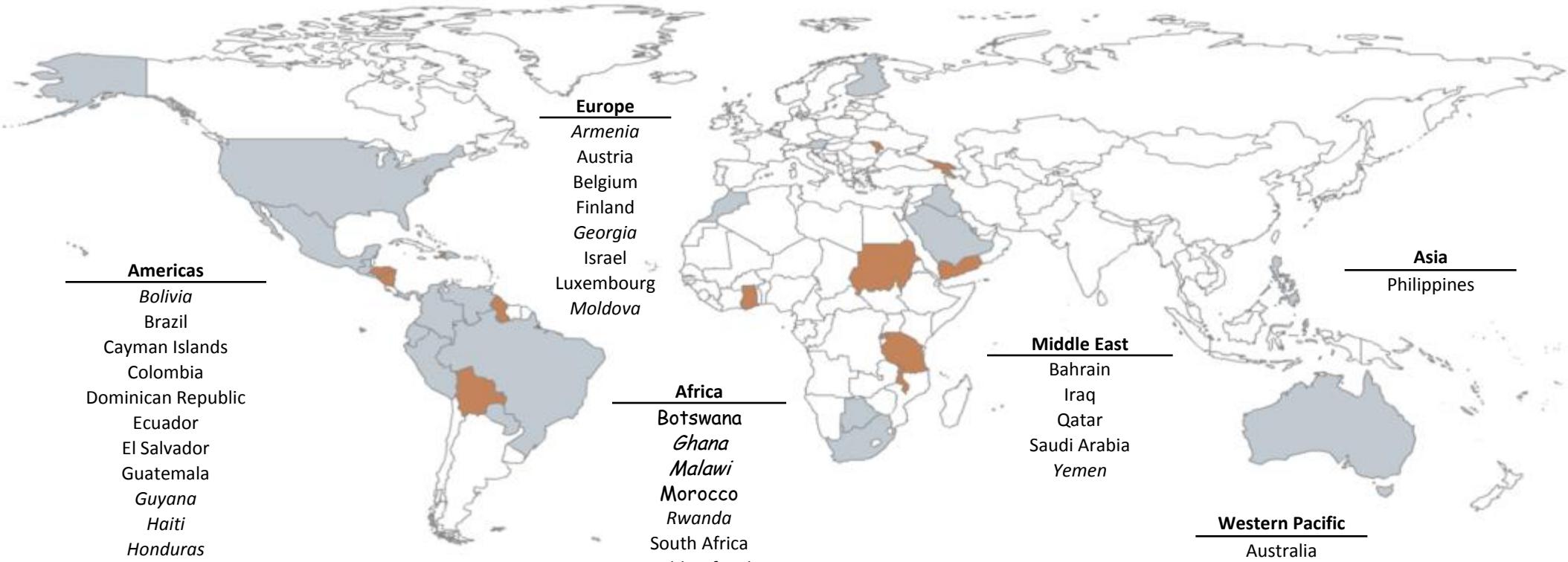


Pentavalent



**Reducción de hospitalización por
GEA por rotavirus**

National RV introductions by geographic region: 45 countries*



*National introductions by geographic region, as of 27 April 2013
RV= rotavirus vaccine



Not GAVI-eligible [31]

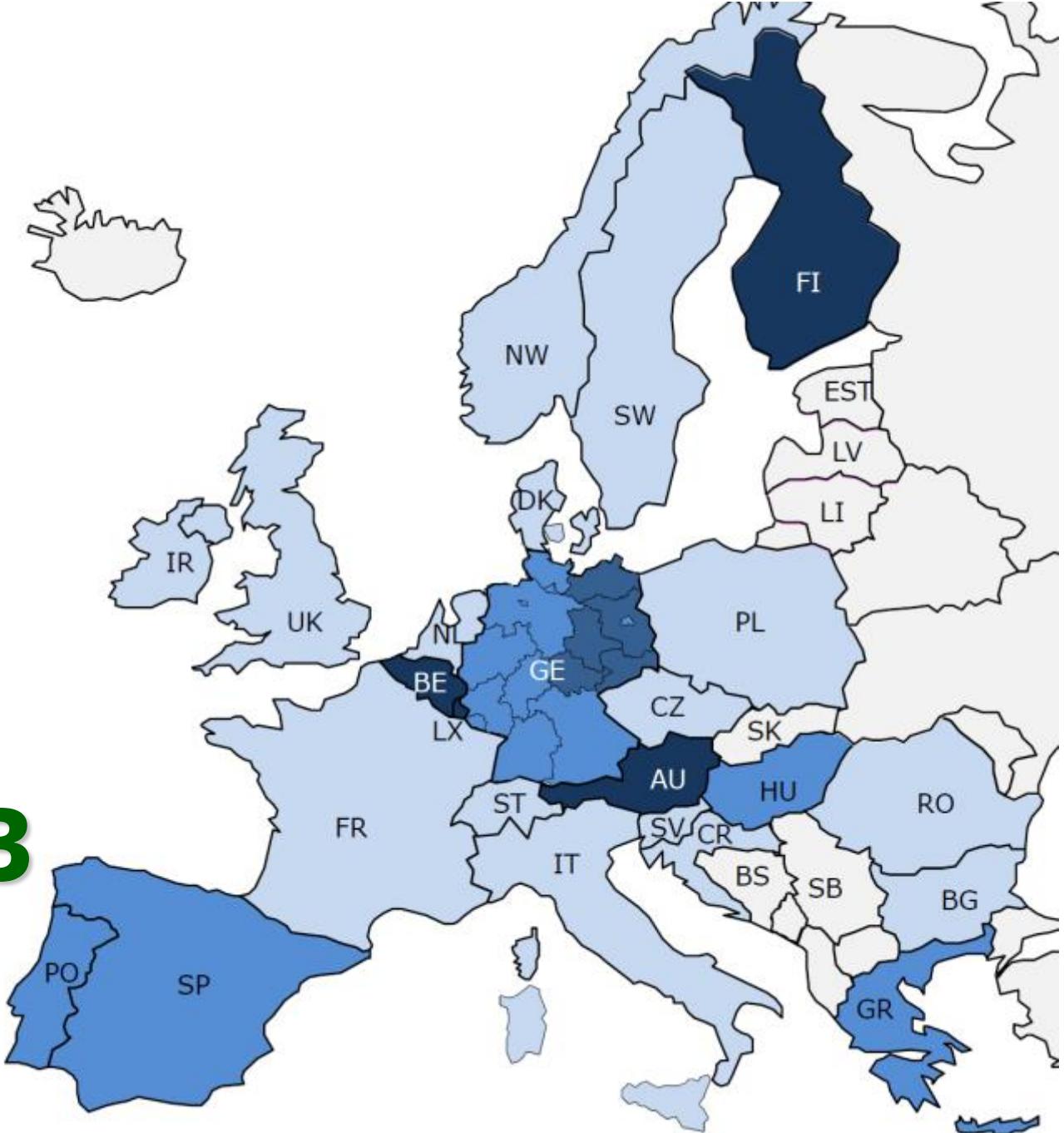


GAVI-eligible [14]



- >90%
- 40–90%
- 20–40%
- <10%
- Not available (NA)

ROTAVIRUS VACCINE COVERAGE EUROPE-2013



(2005-2009)



(2009-to date)



Vaccine Advisory Board Pediatric Spanish Association

- Difusion – Information –
Sensibilization- Education**
 - Professionals**
 - Population**
 - Policy makers / authorities**
- Multidisciplinar Working
groups – CPGs & Consensus**
- Promote active research
(local/national level)**
 - Surveillance**
 - Epidemiological**
 - Pharmacoэкономical**
 - Effectiveness**

THE ROLE OF THE ROTAVIRUS VACCINE IN CHILDHOOD VACCINATION SCHEDULES

F. Giménez Sánchez, F. Martínón Torres, E. Bernaola Iturbe, M. Baca Cots,
F. de Juan Martín, J. Díez Delgado, M. Garcés Sánchez, J.A. Gómez Campderá,
J. Picazo y V. Pineda Solas

An Pediatr (Barc). 2006;64(6):573-7

Impacto familiar de la gastroenteritis por rotavirus en menores de dos años

F. Giménez Sánchez¹, A. Delgado Rubio², F. Martínón Torres³, F. Asensi Botet⁴, M. Miranda Valdivieso⁵, J.L. Gómez 2º apellido¹, S. Alfayate Miguélez⁶, A. Carmona 2º apellido⁷, J. Romero González⁸, M. Crespo Hernández⁹, M. Baca Cots¹⁰, P. Solís Sánchez¹¹, J.A. López Soler¹², M.J. Lozano de la Torre¹³, J. Ruiz 2º apellido¹⁴, V. Pineda Solas¹⁵, I. Manrique Martínez¹⁶, J. García Pérez¹⁷, E. Bernaola Iturbe¹⁸, J.M. del Valle Millán¹⁹, M. Moya Benavent²⁰, L. Ortigosa Castillo²¹, I. Romero Blanco²², E. Román Riechmann²³, J.M. Vizcay Vilella²⁴, M.A. Sesmero Lillo²⁵, C. Rodrigo Gonzalo de Liria²⁶, E. Serrano Poveda²⁷, F. de Juan Martín²⁸ y M. Loriente Tur²⁹, en representación del Grupo ROTASCORE

An Pediatr (Barc). 2008;69(6):515-20

Acta Pædiatrica ISSN 0803-5253 ©2010

REGULAR ARTICLE

Multicenter prospective study analysing the role of rotavirus on acute gastroenteritis in Spain

■ F Giménez-Sánchez (dr.gimenez@cajamar.es)¹, A Delgado-Rubio², F Martínón-Torres³, E Bernaola-Iturbe⁴, Rotascore Research Group*

1.Pediatric Infectious Diseases Unit, Hospital Torrecardenas, Almería, Spain

2.Pediatrics Department Hospital de Madrid, Madrid, Spain

3.Hospital Clínico Universitario de Santiago, Santiago de Compostela, Spain

4.Hospital Virgen del Camino, Pamplona, Spain

Results: A total of 1192 children were enrolled (mean age: 11.2 months). Fever, Vomiting, weakness and dehydration were more frequent in rotavirus-positive AGE cases. Severity score was

An estimation of indirect costs caused by acute rotavirus gastroenteritis in a Galician area, Spain

Federico Martínón-Torres • Marta Bouzón-Alejandro •
Maria López-Sousa • Lorenzo Redondo-Collazo •
Santiago Almeida-Agudín •
Consuelo Astorgano-Fernández •
José María Martínón-Sánchez

Eur J Pediatr. 2008 Mar;167(3):337-9 [Epub 2007 Mar 16]

Proyecto ROTACOST www.rotacost.org



Bouzón-Alejandro et al. BMC Pediatrics 2011, 11:81
<http://www.biomedcentral.com/1471-2431/11/81>

RESEARCH ARTICLE

Open Access

Prospective evaluation of indirect costs due to acute rotavirus gastroenteritis in Spain: the ROTACOST study

Marta Bouzón-Alejandro^{1,3†}, Lorenzo Redondo-Collazo^{1,2,3}, Juan Manuel Sánchez-Lastres^{3,4}, Nazareth Martínón-Torres^{1,2,3}, José María Martínón-Sánchez^{1,2,3} and Federico Martínón-Torres^{1,2,3*†}, for the ROTACOST research team⁵

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favoritos

Multiple publications
Formation activities
Clinical guides



Martínón-Torres, ESPID 2012

Vacunación frente a **ROTAVIRUS**

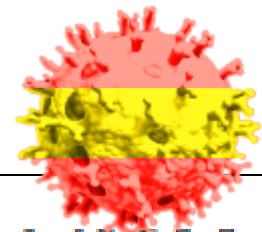
Documento de Consenso
de las Sociedades Científicas

6 de Marzo de 2008



Sociedad Española de Urgencias de Pediatría

- **DR. DÍEZ DOMINGO, JAVIER**
DIRECTOR ÁREA DE VACUNAS
C. SUPERIOR DE INVEST. SALUD PÚBLICA (CSISP). VALENCIA
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HOSPITAL SAN JUAN DE DEU. BARCELONA
- **DR. GIMÉNEZ SÁNCHEZ, FRANCISCO**
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CLÍNICA MEDITERRÁNEO. ALMERÍA
- **DR. MARTINÓN TORRES, FEDERICO**
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COMPL. HOSP. UNIV. SANTIAGO. SANTIAGO
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- **DRA. WILHELMÍ DE CAL, ISABEL**
JEFE DE MICROBIOLOGÍA
HOSPITAL SEVERO OCHOA. MADRID



IMMUNIZATION SCHEDULE OF THE SPANISH ASSOCIATION OF PEDIATRICS: RECOMMENDATIONS 2008

E. Bernaola Iturbe, F. Giménez Sánchez, M. Baca Cots, F. de Juan Martín, J. Díez Domingo, M. Garcés Sánchez, A. Gómez-Campderá, F. Martinón Torres, J.J. Picazo y V. Pineda Solás

Comité Asesor de Vacunas de la Asociación Española de Pediatría.

Varicela ¹¹					Var		Var	Varicela
Neumococo ¹²		Pn7v	Pn7v	Pn7v	Pn7v			
Papilomavirus ¹³								VPH
Rotavirus ¹⁴⁻¹⁵			ROTAV					

Effectiveness of rotavirus vaccination in Spain

Federico Martínón-Torres,^{1-3,†,*} Marta Bouzón Alejandro,^{1,3,†} Lorenzo Redondo Collazo,¹⁻³ Juan Manuel Sánchez Lastres,^{3,4} Sonia Pértiga Díaz,⁵ M^a Teresa Seoane Pillado,⁵ José María Martínón Sánchez¹⁻³ and ROTACOST research team⁶

- October 2008-June 2009
- 682 children below 5 yo with AGE prospectively collected
- 18 C.S. y 10 hospitals from Galicia and Asturias
- In all cases, rotavirus antigen detection was performed (Test Vikia®)
- Case-control design VE= 1-OR

Vaccine effectiveness to prevent any rotavirus AGE

91.5%

(CI 95%:83.7%-95.6%)

Complete vacc: **92.8%** (84.7-96.6%)

Partial vacc: **84.0%** (45.5-95.3%)

Vaccine effectiveness to prevent hospital admission

95.6%

(CI 95%: 85.6-98.6%)

Complete vacc: **98.3%** (87.4-100%)

Partial vacc: **89.4%** (53.9-97.5%)



Effectiveness of rotavirus vaccines in preventing cases and hospitalizations due to rotavirus gastroenteritis in Navarre, Spain

Jesús Castilla^{a,b,*}, Xabier Beristain^c, Víctor Martínez-Artola^c, Ana Navascués^d, Manuel García Cenoz^{a,b}, Nerea Álvarez^e, Isabel Polo^c, Ana Mazón^c, Alberto Gil-Setas^c, Aurelio Barricarte^{a,b}

Table 3

Rotavirus vaccine effectiveness in preventing hospital admission due to rotavirus gastroenteritis in children aged 3–59 months.

	Cases/controls	Crude OR (95% CI)	Adjusted OR (95% CI) ^a	Vaccine effectiveness (95% CI)
Total				
Unvaccinated	249/390	1	1	
Fully vaccinated	9/80	0.18 (0.09–0.36)	0.17 (0.07–0.35)	83% (65–93%)
Aged <24 months				
Unvaccinated	207/302	1		
Fully vaccinated	8/69	0.17 (0.08–0.36)	0.18 (0.07–0.40) ^b	82% (60–93%) ^b
Aged ≥24 months				
Unvaccinated	42/88	1	1	
Fully vaccinated	1/11	0.19 (0.02–1.52)	0.11 (0.002–0.83) ^b	89% (17–99.8%) ^b
Period 2008–2009				
Unvaccinated	91/237	1	1	
Fully vaccinated	7/29	0.63 (0.27–1.49)	0.61 (0.21–1.54) ^c	39% (−54 to 79%) ^c
Period January 2010–June 2011				
Unvaccinated	158/153	1	1	
Fully vaccinated	2/51	0.04 (0.01–0.16)	0.04 (0.004–0.15) ^c	96% (85–99%) ^c

^a Odds ratio and 95% confidence interval obtained by exact logistic regression adjusted for age group, birth year and major chronic conditions.

^b Comparison of vaccine effectiveness in preventing cases with age <24 months vs ≥24 months: $p = 0.9601$.

^c Comparison of vaccine effectiveness in the period 2008–2009 vs the period January 2010–June 2011: $p = 0.0009$.

Efectividad de la vacunación frente al rotavirus en Valencia

Figure 1. Total number of children by year of birth and vaccination status

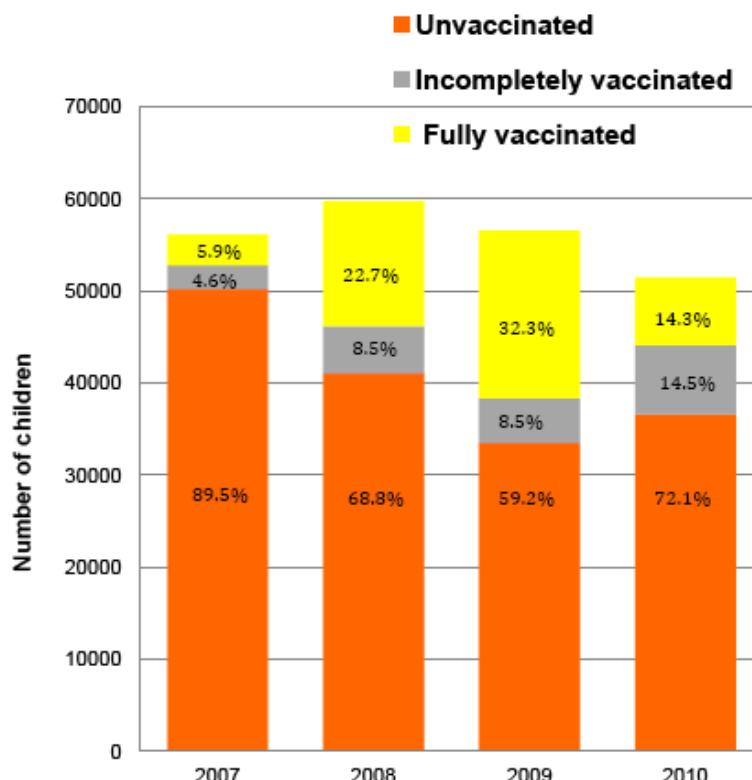


Figure 2. Hospitalizations for rotavirus AG

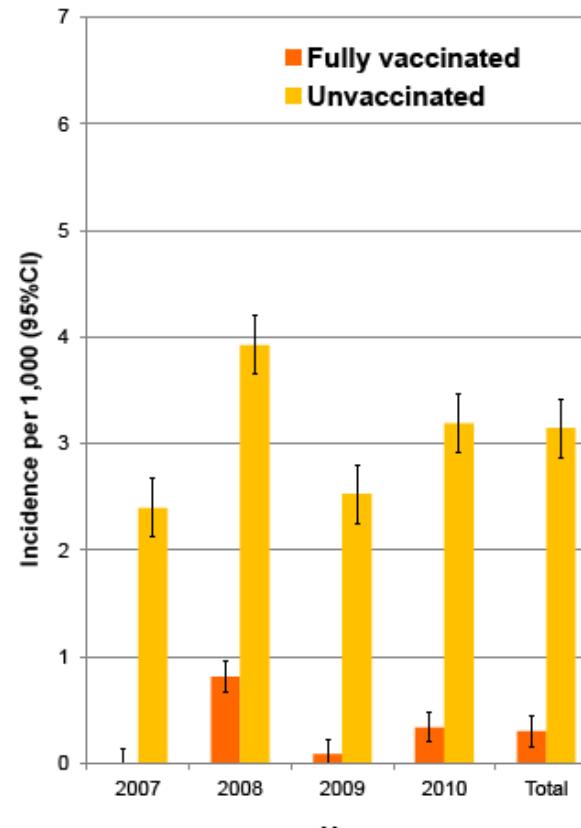
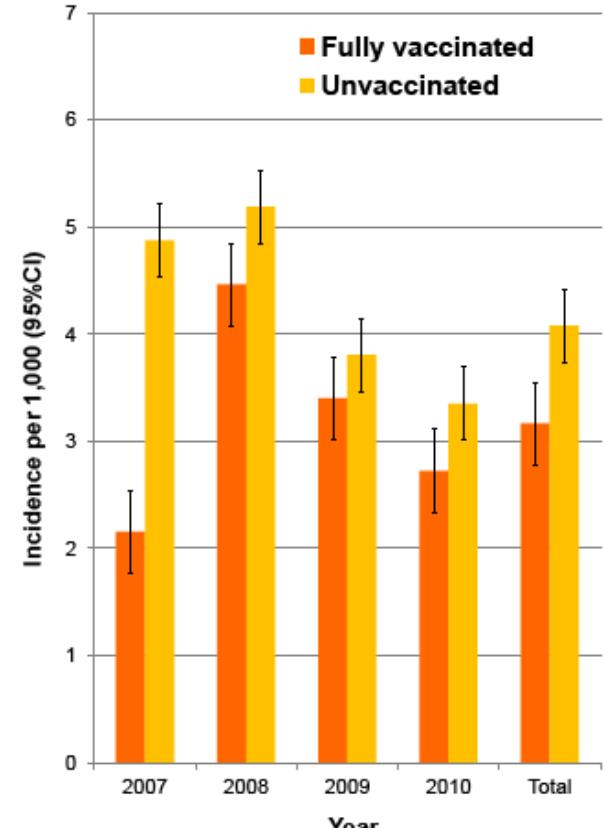


Figure 3. Hospitalizations for other AG

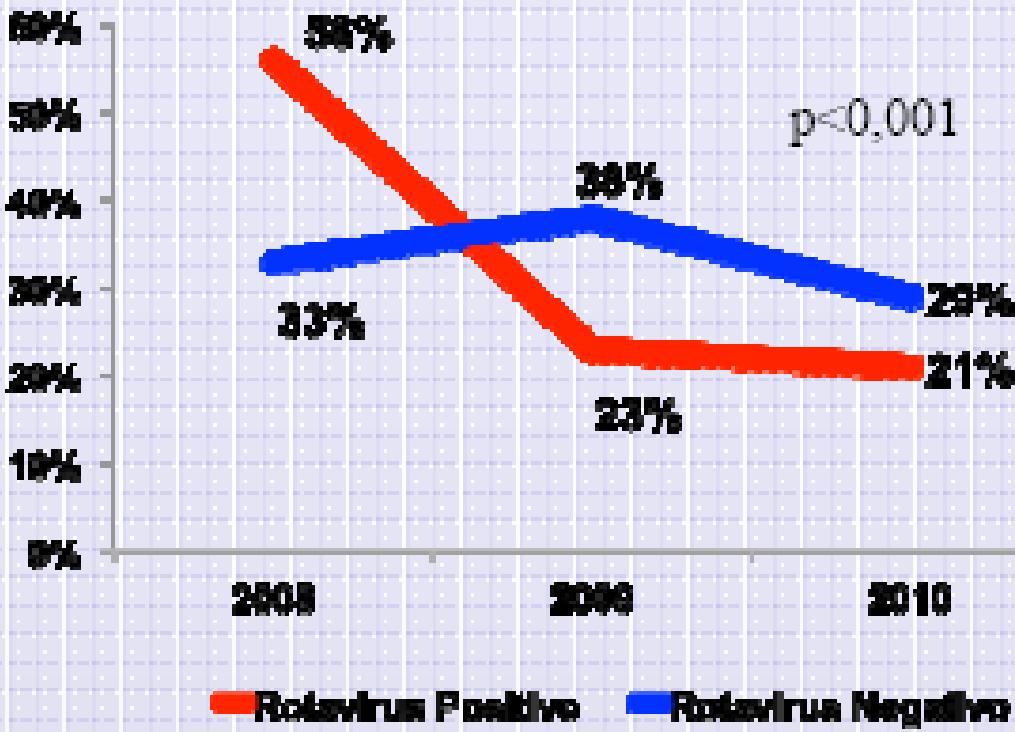


Efectividad RV hospit **90.4% (85-94)**

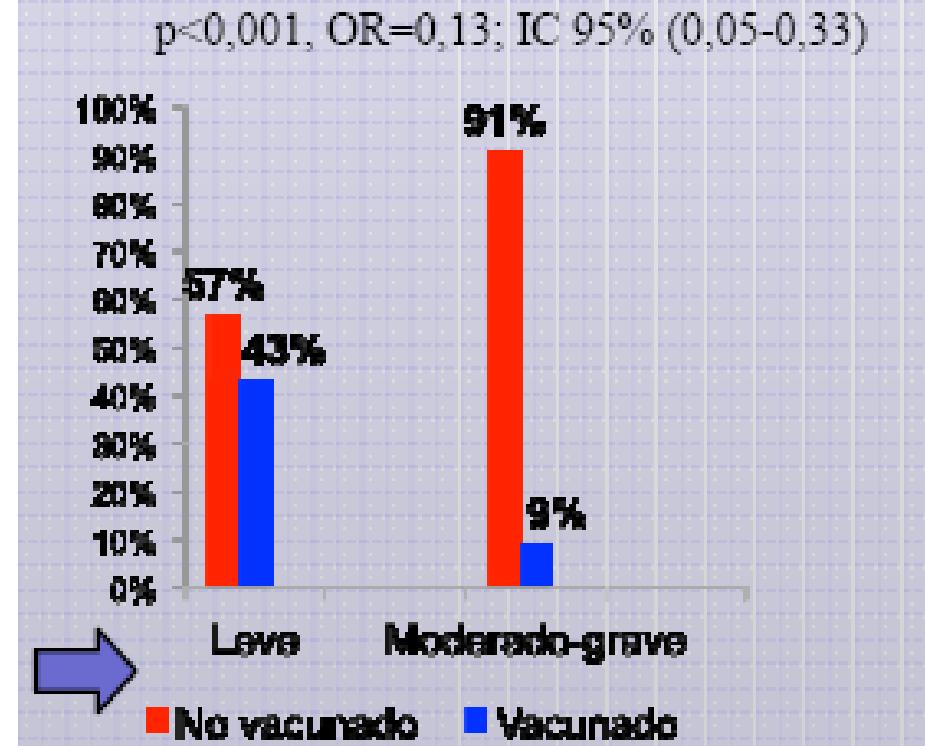
Diez Domingo J. et al. ESPID 2012

Efectividad de la vacunación frente al rotavirus en Almería

Evolución de las GEAs según años



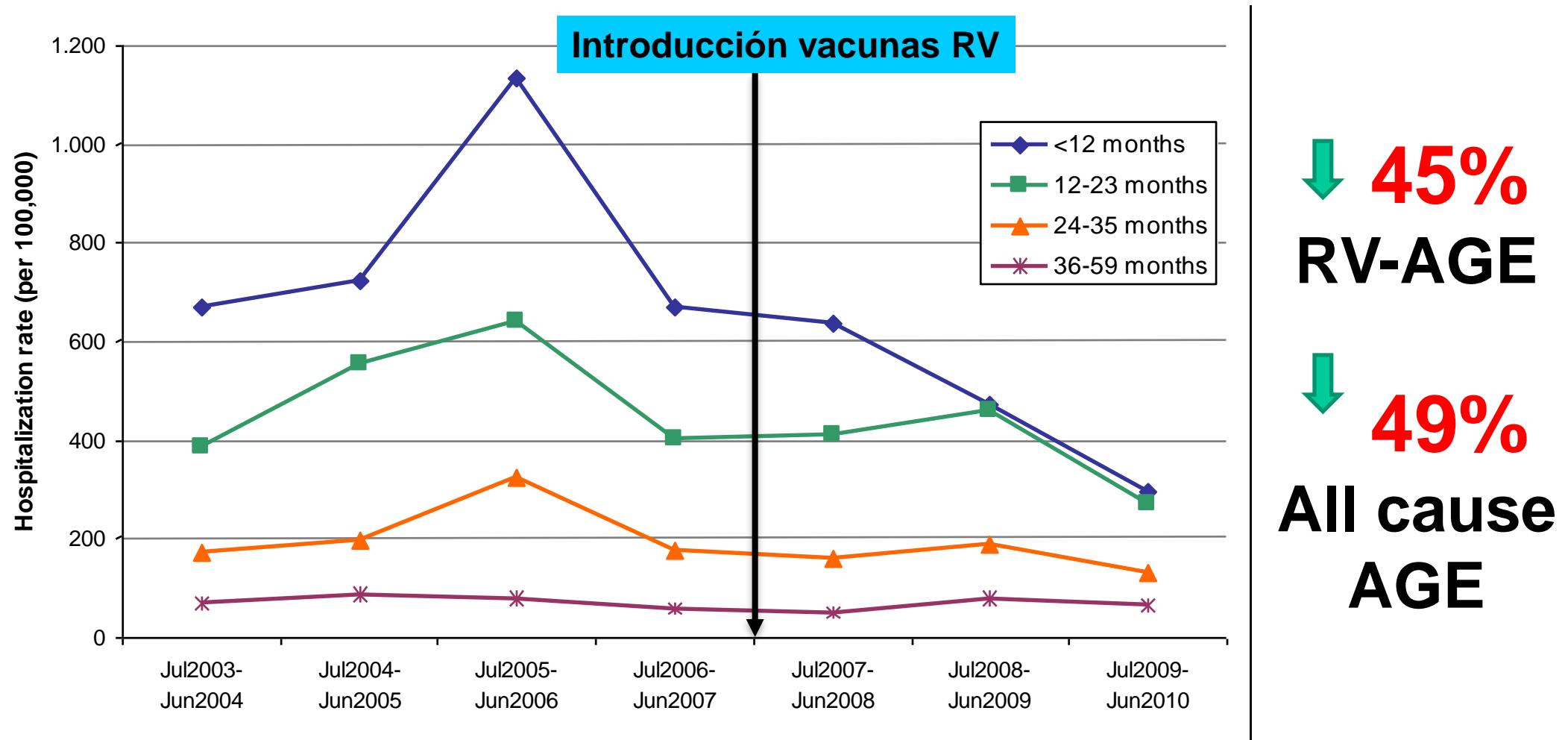
Gravedad clínica (Score Vesikari)



Efectividad RV hospital 90.4% (85-94)

Cobos E, et al. SEIP 2012

Impacto de la vacunación frente a rotavirus en Galicia



Martinón-Torres F, et al. Human Vaccine & Immuno, vol 8, issue 7, July 2012, epub ahead of print



Impact of rotavirus vaccination on hospitalizations for rotavirus diarrhea: The IVANHOE study

Arnaud Gagneur ^{a,e,*}, Emmanuel Nowak ^b, Thomas Lemaitre ^b, Jean-Francois Segura ^a, Nadège Delaperrière ^a, Lydie Abalea ^a, Elise Poulhazan ^b, Anne Jossens ^a, Lucie Auzanneau ^b, Adissa Tran ^c, Christopher Payan ^c, Nadine Jay ^a, Loïc de Parscau ^a, Emmanuel Oger ^d, The IVANHOE investigators¹

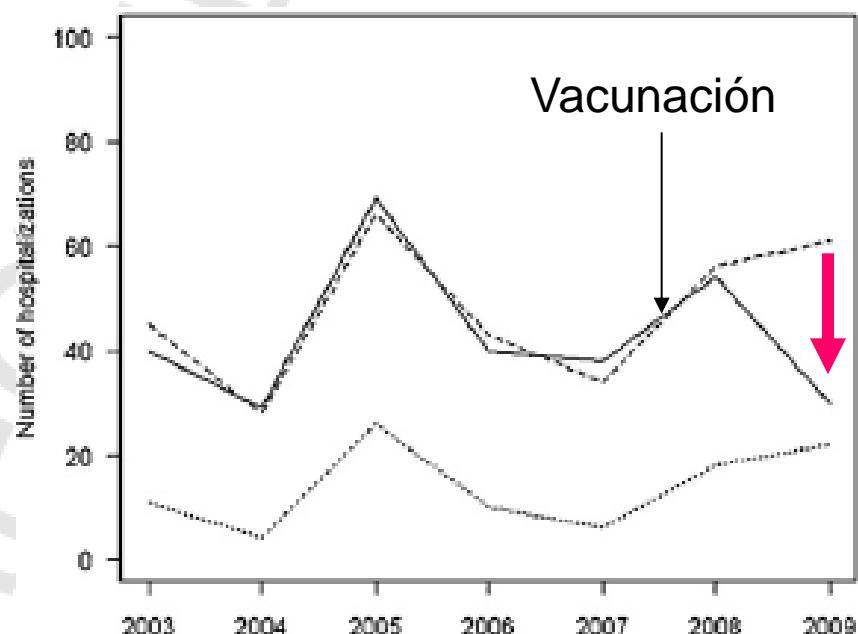


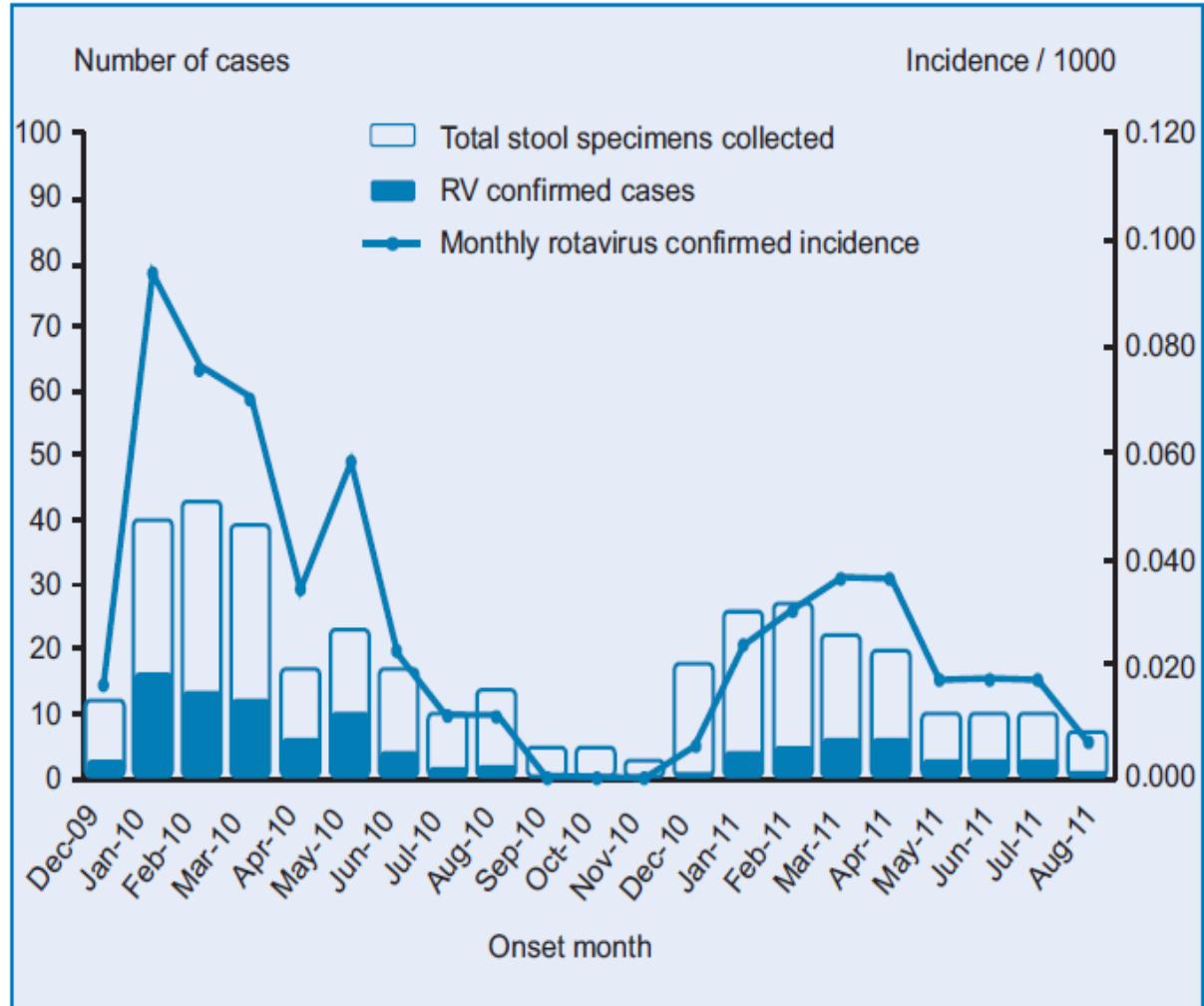
Fig. 4. Observed (<2 and 2–5 years old; continuous and dotted lines) and expected (<2 years old; dashed line) numbers of hospitalizations during epidemic seasons.

Prospectivo – Pentavalente
Vigilancia 5 años previos + 2
Cobertura: **47.1%**
Tiene en cuenta las variaciones estacionales e interpeidémicas
RRR hospitalización = **98% (83-100%)**

FINLANDIA – 2 AÑOS TRAS INICIO

VACUNACIÓN UNIVERSAL ROTAVIRUS

Figure 4. Total number of stool specimens collected, number of RV confirmed cases and RV confirmed incidence by month of AGE onset, overall and by study site, Tampere and Oulu, Finland, 2009-2011

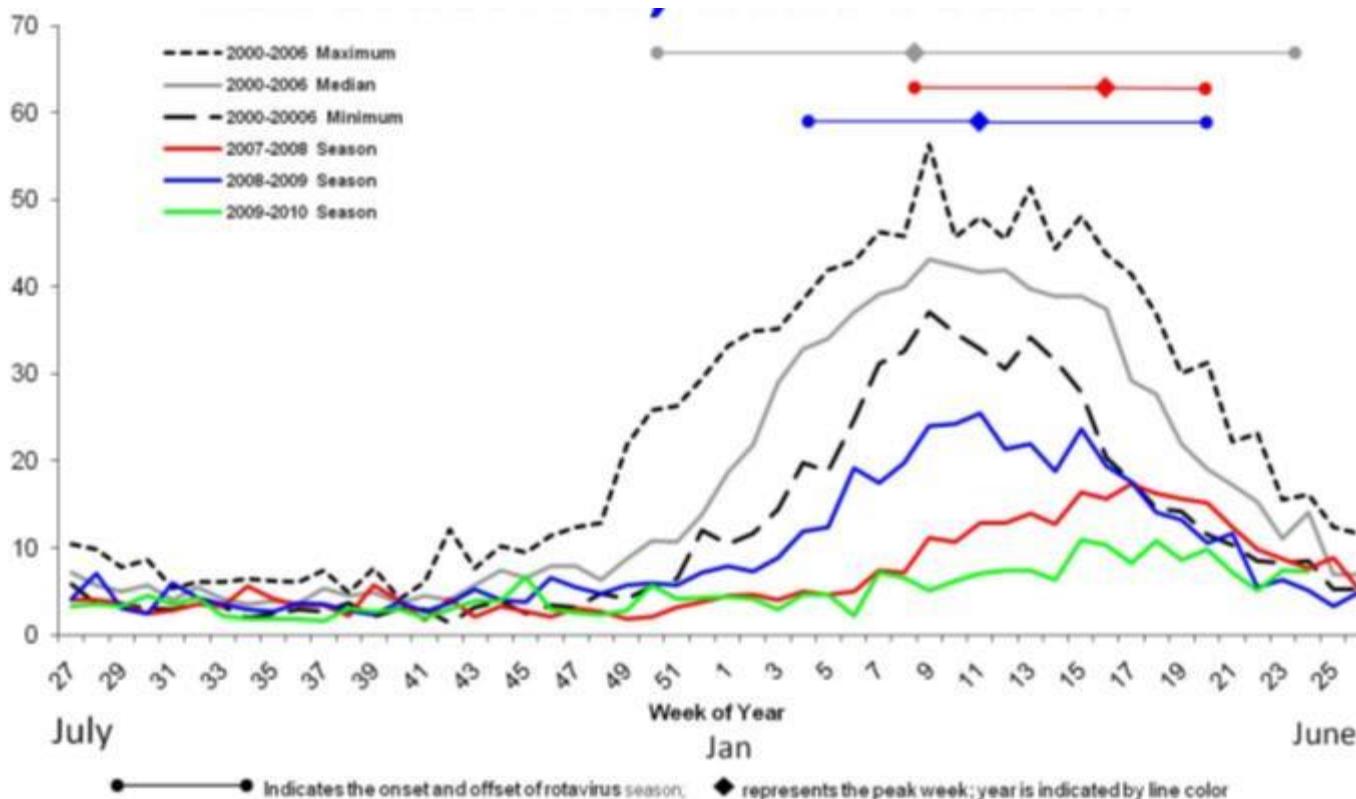


- Pentavalent vaccine included in Finnish National Vaccination Program in September 2009 (schedule 2/3/5 months)
- Vaccine coverage rapidly reached approximately 95-97%
- As of 08-2011, 301 AGE (100 RVGE)
- Vaccine effectiveness estimated > 95% in children born after December 2007

Vesikari T et al. ESPID 2012 - Poster nº 683
Vesikari T et al. Eur J Ped 2013 epub ahead

Sustained Decline in Rotavirus Detections in the United States Following the Introduction of Rotavirus Vaccine in 2006

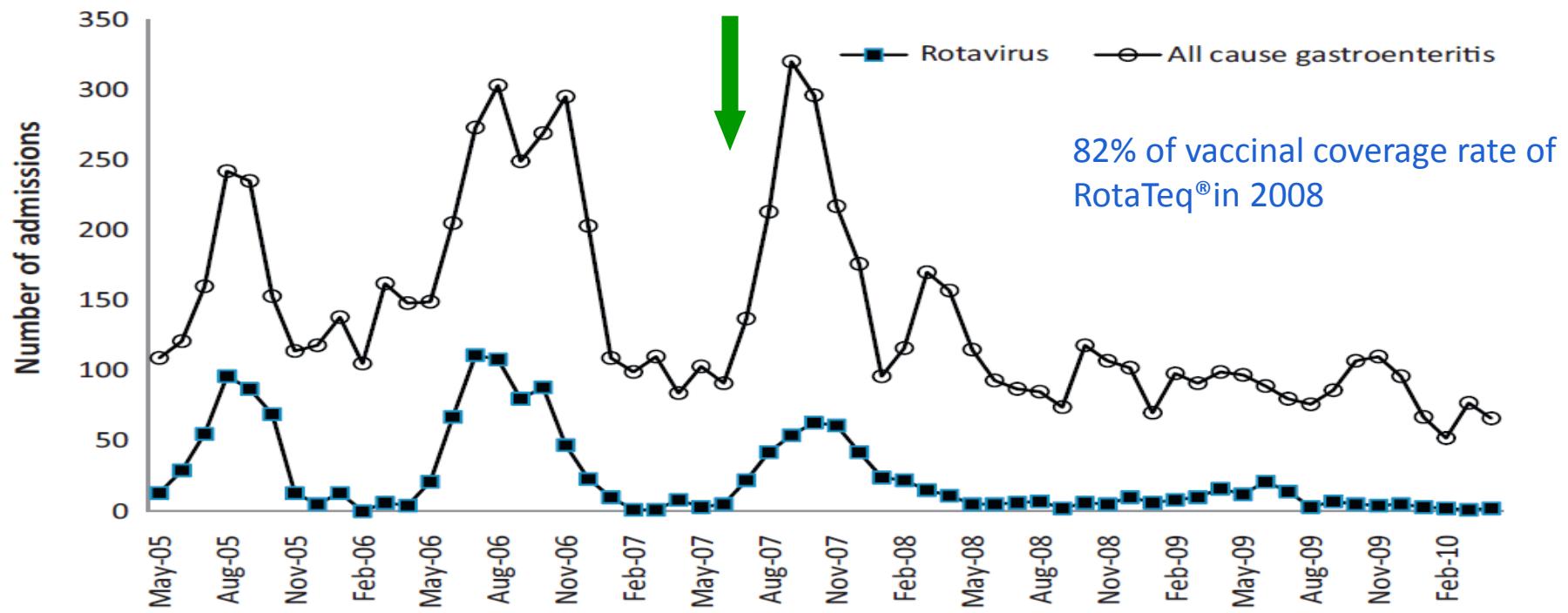
Jacqueline E. Tate, PhD, Jeffry D. Mutuc, MPH, Catherine A. Panozzo, MPH,
Daniel C. Payne, PhD, Margaret M. Cortese, MD, Jennifer E. Cortes, MD, Catherine Yen, MD, MPH,
Douglas H. Esposito, MD, Benjamin A. Lopman, PhD, Manish M. Patel, MSc, MD,
and Umesh D. Parashar, MB BS, MPH



- Cobertura (≥ 1 d): 72 % cobertura vacunal (al menos 1 dosis)
- Actividad rotavirus disminuida y retrasada
- **NO** cumplió criterios de inicio de temporada!!

South Australia - 3 years after pentavalent rotavirus vaccine introduction

Introduction of RotaTeq® in children
vaccination schedule (2/4/6) in July 2007



Gastroenteritis Hospital admissions in children < 6 years of age linked to RV or all causes, per month.

Retrospective analysis of hospitals admissions conducted in children < 6 years of age in South Australia

JAMA. 2009;301(21):2243-2251 | Patel; Cristina Pedreira; Lucia Helena De Oliveira; et al.

Vesikari severity score ≥11

Cases 110/146 (75)

Controls

Neighborhood 338/396 (85) 0.48 (0.28 to 0.82) 52 (18 to 72)

Hospital 278/321 (87) 0.37 (0.20 to 0.66) 63 (34 to 80)

Combined 616/717 (86) 0.42 (0.26 to 0.70) 58 (30 to 74)

Vesikari severity score ≥15

Cases 28/39 (72)

Controls

Neighborhood 86/99 (87) 0.27 (0.09 to 0.82) 73 (18 to 91)

Hospital 81/93 (87) 0.14 (0.04 to 0.55) 86 (46 to 96)

Combined 167/192 (87) 0.23 (0.08 to 0.61) 77 (39 to 92)

Eficacia frente a hospitalización o visitas a urgencias**72 al 92 %**

Effect of Rotavirus Vaccination on Death from Childhood Diarrhea in Mexico



Vesta Richardson, M.D., Joselito Hernandez-Pichardo, M.D.,
Manjari Quintanar-Solares, M.D., Marcelino Esparza-Aguilar, M.D.,
Brian Johnson, B.S., Cesar Misael Gomez-Altamirano, M.D.,

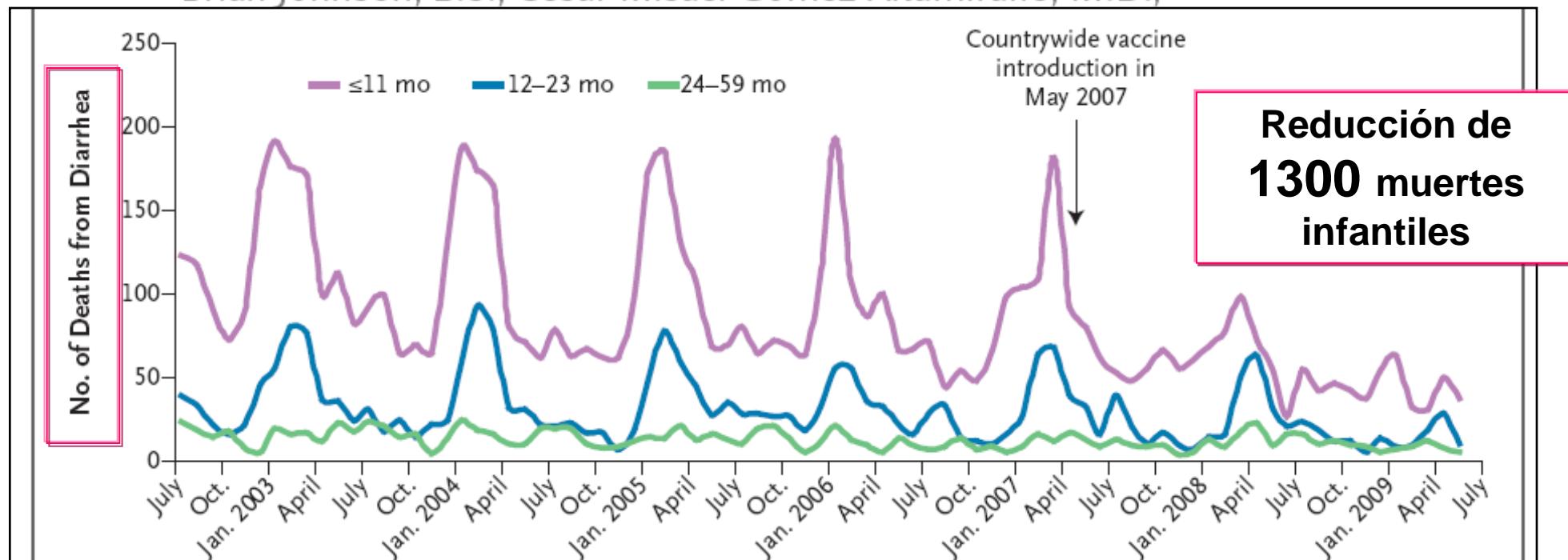
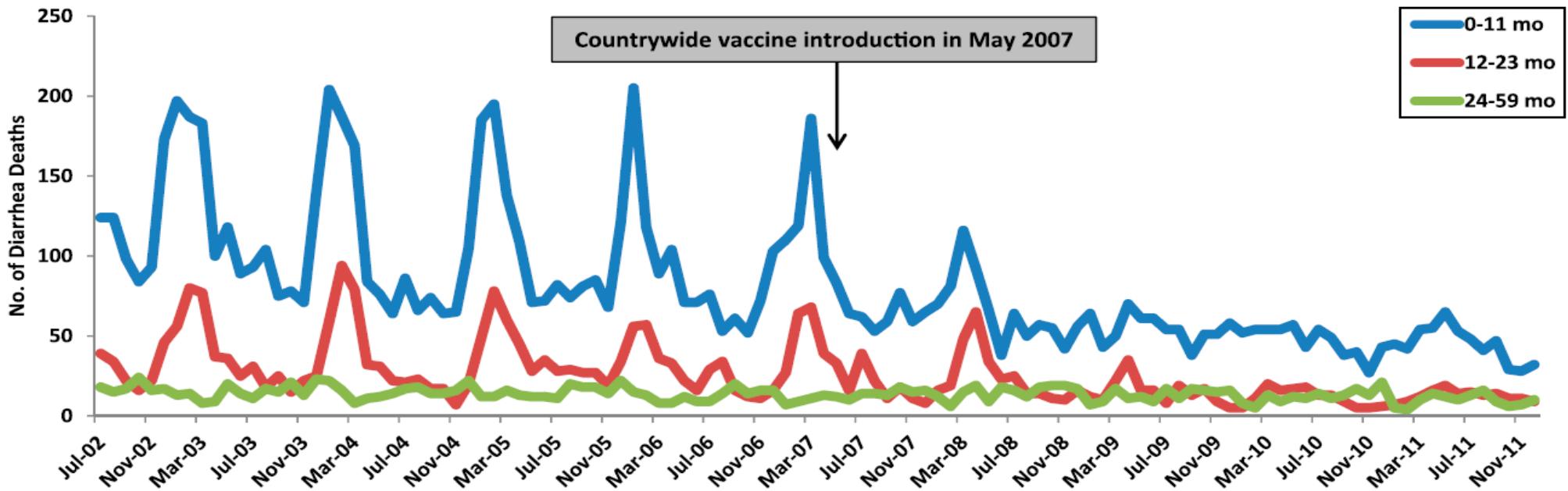


Figure 1. Number of Diarrhea-Related Deaths among Children 59 Months of Age or Younger from July 2002 through May 2009 in Mexico, According to Age Group.

EFFECT OF ROTAVIRUS VACCINATION ON ACUTE DIARRHEA DEATHS IN MEXICAN CHILDREN



Age group (months)	2003 -2006 period		Post-vaccine		Absolute reduction		Relative reduction		
	Median	Diarrhea-Related Rate of Death	Median	Diarrhea-Related Rate of Death	No.	%	% of reduction	IC 95%	P
0 to 59	1793	16.87	872	9.14	921	7.72	45.8	39.8 – 51.7	<0.001
0 to 11	1197	59.65	601	31.54	596	28.1	47.1	40.0 - 54.2	<0.001
12 to 23	421	19.53	155	8.13	266	11.41	58.4	46.5 – 70.3	<0.001
24 to 59	175	2.71	116	2.03	59	0.68	25.1	4.8 – 45.4	0.015



GREEN APPLES – CUESTIONES ABIERTAS

- 1.- Lactancia materna y prematuros**
- 2.- ¿Cuánto dura la protección?**
- 3.- Impacto epidemiológico**
- 4.- Inmunidad Colectiva**
- 5.- Otras formas de infección**

Safety and Efficacy of the Pentavalent Human-Bovine (WC3) Reassortant Rotavirus Vaccine in Healthy Premature Infants

Michelle G. Goveia, MD,* Zoe M. Rodriguez, MD,† Michael J. Dallas, PhD,* Robbin F. Itzler, PhD,* John W. Boslego, MD,* Penny M. Heaton, MD,* and Mark J. DiNubile, MD*
for the REST Study Team

The Pediatric Infectious Disease Journal • Volume 26, Number 12, December 2007



TABLE 4. Rate Reduction for Hospitalizations and Emergency Department Visits Attributable to Rotavirus Gastroenteritis Regardless of Serotype in Premature Infants

	Vaccine	Placebo	Rate Reduction (%)	95% CI
Per-protocol population starting 2 weeks after the third dose				
Number of evaluable premature infants completing the full 3-dose series	764	818		
Number of hospitalizations and emergency department visits	0	21	100	82.2–100
Number of hospitalizations	0	9*	100	53.0–100
Number of emergency department visits	0	12†	100	66.6–100
Intention-to-treat population anytime after the first dose				
Number of evaluable premature infants receiving at least 1 dose	938	990		
Number of hospitalizations and emergency department visits	1	24	95.5	76.4–99.9
Number of hospitalizations	0	10‡	100	58.8–100
Number of emergency department visits	1§	14	92.3	56.6–99.8

N= 2070 pacientes

Edad 25-36 sem [34 s]

NO efectos adversos

↓↓↓ 100% (82-100)

Hospitalización y
Asistencia a Urgencias

Rotavirus Vaccination of Very Low Birth Weight Infants at Discharge From the NICU



WHAT'S KNOWN ON THIS SUBJECT: Preterm and low birth weight infants are at increased risk of hospitalization due to rotavirus gastroenteritis, and rotavirus vaccine is immunogenic and well tolerated among these infants when provided at or after discharge from the NICU.



WHAT THIS STUDY ADDS: Many preterm infants with a birth weight of ≤ 1500 g are not eligible to receive rotavirus vaccination because they remain in the NICU beyond the upper age limit recommended for immunization. New strategies are needed.

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KEY WORDS

rotavirus, immunization, preterm infant, neonatal ICU

ABBREVIATIONS

AAP—American Academy of Pediatrics

ELBW—extremely low birth weight

PMH—Parkland Memorial Hospital

RV5—RotaTeq

VLBW—very low birth weight

Por edad insuficiente, excesiva u omisión,
la mayoría acaban NO vacunándose

EFFICACY OF PENTAVALENT HUMAN-BOVINE (WC3) REASSORTANT ROTAVIRUS VACCINE BASED ON BREASTFEEDING FREQUENCY



Michelle G. Goveia, MD, MPH, Mark J. DiNubile, MD,
Michael J. Dallas, PhD, Penny M. Heaton, MD, and
Barbara J. Kuter, PhD, MPH; for the REST Study Team



The Pediatric Infectious Disease Journal • Volume 27, Number 7, July 2008

NO interferencia de la lactancia materna (exclusiva o mixta)

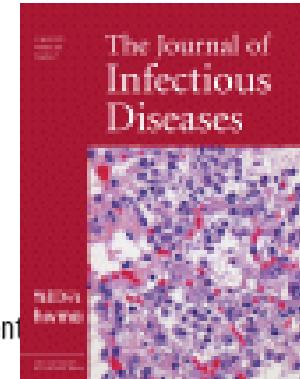
TABLE 1. Efficacy Against Any and Severe G1-4 Rotavirus Gastroenteritis Through the First Full Rotavirus Season Based on Frequency of Breastfeeding (Per Protocol Population and Analysis)

Breastfeeding Frequency	Any G1-4 Rotavirus Gastroenteritis			Severe G1-4 Rotavirus Gastroenteritis		
	No. Cases/Number Vaccinated		Efficacy (95% CI)	No. Cases/Number Vaccinated		Efficacy (95% CI)
	Vaccine Recipients	Placebo Recipients		Vaccine Recipients	Placebo Recipients	
Never	19/815	60/817	68.3% (46.1–82.1)	0/815	9/817	100% (48.3–100)
Some	24/953	133/947	82.2% (72.3–89.0)	1/953	22/947	95.4% (71.3–99.9)
Exclusively	39/767	122/799	68.0% (53.8–78.3)	0/767	20/799	100% (79.3–100)

(Rotarix) against Severe Diarrhea Caused by Serotypically Unrelated G2P[4] Strains in Brazil

Jailson B. Correia,¹ Manish M. Patel,² Osamu Nakagomi,^{3,4} Fernanda M. U. Montenegro,¹ Eliane M. Germano,¹ Nancy B. Correia,¹ Luis E. Cuevas,⁵ Umesh D. Parashar,² Nigel A. Cunliffe,⁴ and Toyoko Nakagomi^{3,4}

¹Instituto de Medicina Integral Prof Fernando Figueira, Recife, Brazil; ²Centers for Disease Control and Prevention, Atlanta, Georgia; ³Department of Molecular Microbiology and Immunology, and Global Centre of Excellence, Nagasaki University, Nagasaki, Japan; ⁴Department of Medical Microbiology, University of Liverpool, and ⁵Liverpool School of Tropical Medicine, Liverpool, United Kingdom



Group	G2P[4] rotavirus strains		All rotavirus strains	
	Adjusted OR (95% CI)	Effectiveness (95% CI)	Adjusted OR (95% CI)	Effectiveness (95% CI)
Severe rotavirus disease (requiring hospital admission or treatment at emergency department)				
Case patients 6–11 months of age				
Rotavirus-negative control participants	0.23 (0.09–0.58)	77 (42–91)	0.27 (0.11–0.64)	73 (36–89)
Control participants with ARI	0.23 (0.10–0.57)	77 (43–90)	0.27 (0.12–0.61)	73 (38–88)
Case patients >12 months of age				
Rotavirus-negative control participants	1.24 (0.53–2.90)	–24 (–190 to 47)	1.51 (0.67–3.48)	–51 (–248 to 34)
Control participants with ARI	0.84 (0.36–2.01)	15 (–101 to 64)	1.01 (0.43–2.38)	–1 (–138 to 57)
Rotavirus disease requiring hospitalization				
Case patients 6–11 months of age				
Rotavirus-negative control participants	0.15 (0.05–0.46)	85 (54–95)	0.19 (0.07–0.53)	81 (47–93)
Control participants with ARI	0.17 (0.06–0.49)	83 (51–94)	0.20 (0.08–0.52)	80 (48–92)
Case patients >12 months of age				
Rotavirus-negative control participants	0.95 (0.31–2.88)	5 (–187 to 69)	0.95 (0.31–2.88)	5 (–187 to 69)
Control participants with ARI	0.58 (0.19–1.80)	41 (–79 to 81)	0.58 (0.19–1.80)	41 (–79 to 81)

¿Cuánto dura la protección inducida por la vacuna frente a rotavirus?



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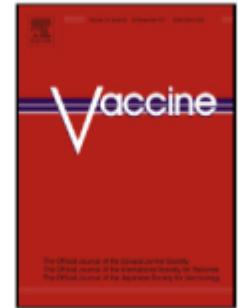
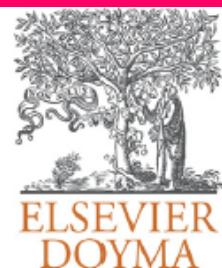


Table 1

Incidence rates per 100,000 in different age groups in the pre-vaccination era 2001–2005 (mean), in 2010 and 2011, and change (in %) in incidence rates in different age groups in the years 2010 and 2011 compared to the mean pre-vaccination era 2001–2005.

	Mean prevaccination era (2001–2005) Incidence rate per 100,000	UMV 2010 Incidence rate per 100,000 (change in %)	UMV 2011 Incidence rate per 100,000 (change in %)
0–<1 year	2141	344 (−84%)	397 (−81%)
1–<2 years	1745	331 (−81%)	332 (−81%)
2–<3.5 years	611	235 (−62%)	216 (−65%)
3.5–<5 years	206	188 (−9%)	201 (−3%)
5–<10 years	53	35 (−34%)	78 (+48%)
10–<15 years	13	9 (−32%)	14 (+6%)
0–<5 years	1008	261 (−74%)	271 (−73%)
0–<15 years	329	98 (−70%)	120 (−64%)

RV1 forma parte del programa de inmunización nacional
La mayoría de los casos son por G2P4



Enfermedades Infecciosas y Microbiología Clínica

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Original breve

Caracterización microbiológica de las gastroenteritis agudas virales atendidas en un servicio de pediatría en un área de alta cobertura vacunal frente a rotavirus

Carlos García-Magán^a, María José de Castro-López^a, José Llovo-Taboada^b, Carmen Curros-Novo^a, Alicia Sánchez-Fauquier^c y Federico Martínón-Torres^{a,*}

^a Departamento de Pediatría, Complejo Hospitalario Universitario de Santiago de Compostela, Santiago de Compostela, España

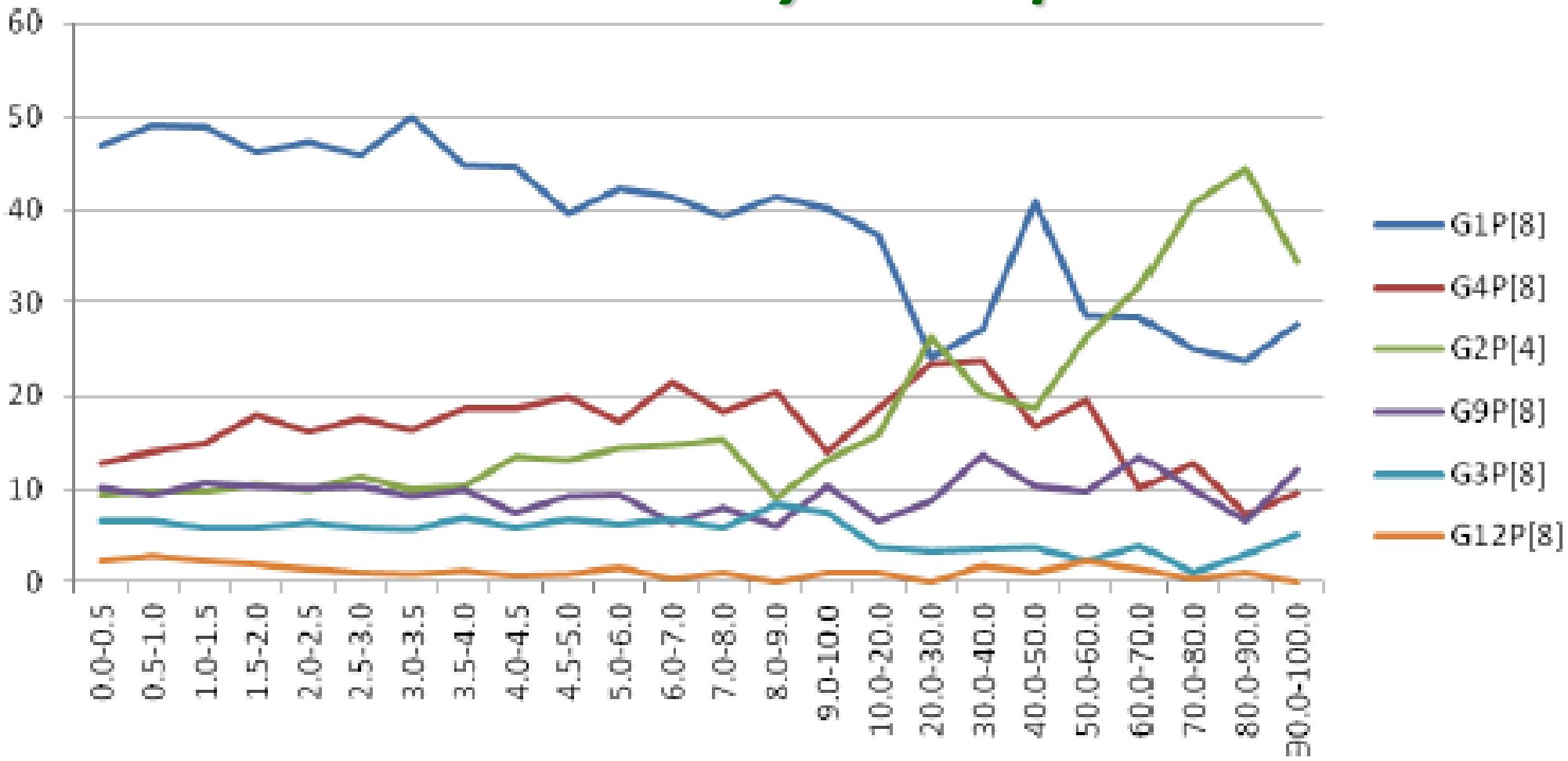
^b Servicio de Microbiología, Complejo Hospitalario Universitario de Santiago de Compostela, Santiago de Compostela, España

^c Instituto de Salud Carlos III, Madrid, España

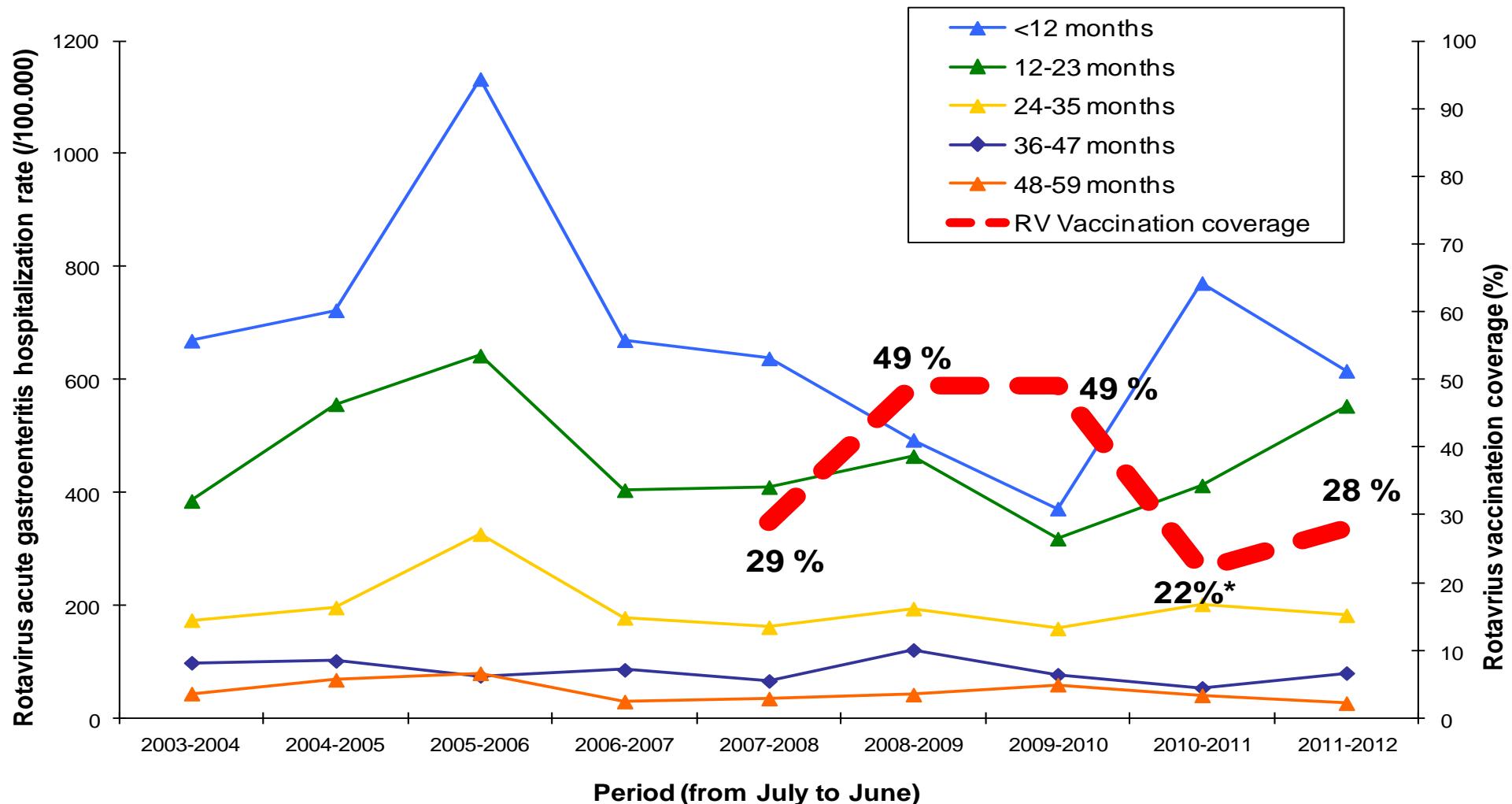
Sin evidencias de reemplazamiento de serotipos en un área con cobertura alta frente a rotavirus

Genotipos rotavirus según edad en EUROPA

EURO-ROTANET 6 years report



Reverse evidence of rotavirus vaccines effectiveness



(*) 22% is the mean RV vaccine coverage for that period. However, for 5 months within that period, no new batches of vaccine were released onto the market, and the coverage estimated for those months was 0-5%.



Jay M. Lieberman^{1,2}, Xiaohua Huang³, Eileen Koski⁴, Harvey Kaufman⁵,
Richard Furlanetto⁶, Herman Hurwitz⁷, Pradeep Ragothaman⁵, Marsha Green⁴

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⁶Quest Diagnostics Nichols Institute, Chantilly, VA; ⁷Quest Diagnostics, Philadelphia, PA

ICAAAC/IDSA 2008

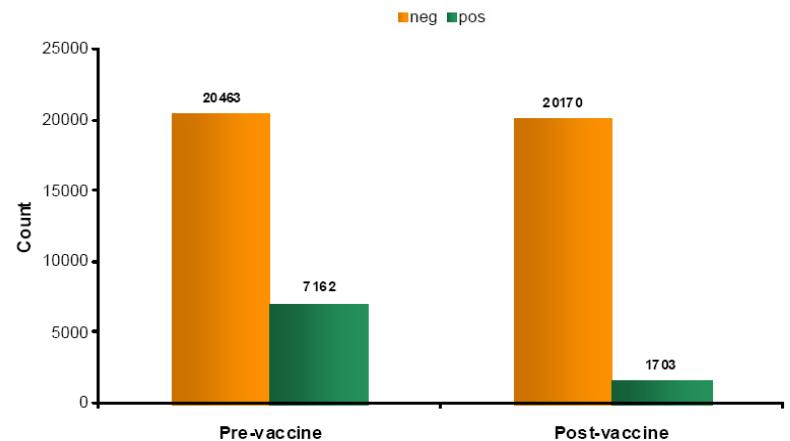


Figure 2. Rotavirus test results before and after vaccine availability.

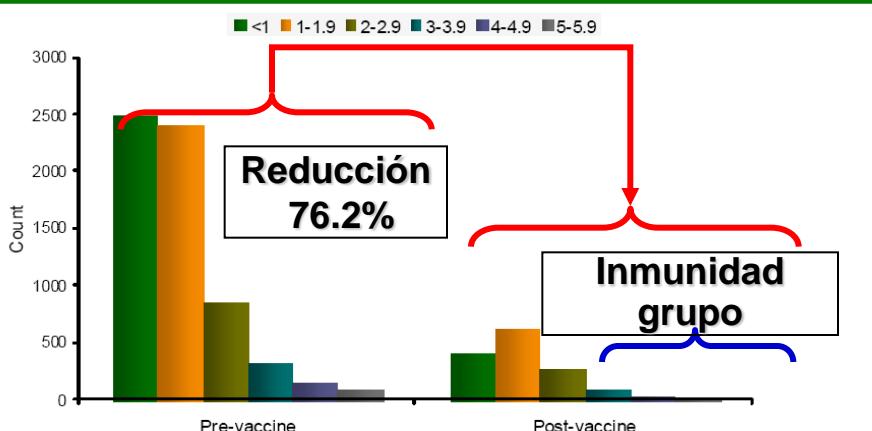


Figure 4. Number of positive rotavirus test results by age (in years), before and after vaccine availability.

- Reducción **76.2%**, a expensas sobre todo de menores de 1 año
- Descenso >>> lógico esperable : Cobertura **49%** con \geq 1 dosis)
- **PROTECCION INDIRECTA**

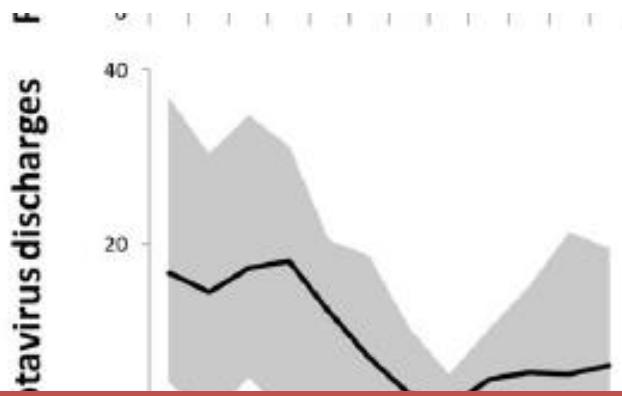
Infant Rotavirus Vaccination May Provide Indirect Protection to Older Children and Adults in the United States

Ben A. Lopman, Aaron T. Curns, Catherine Yen, and Umesh D. Parashar

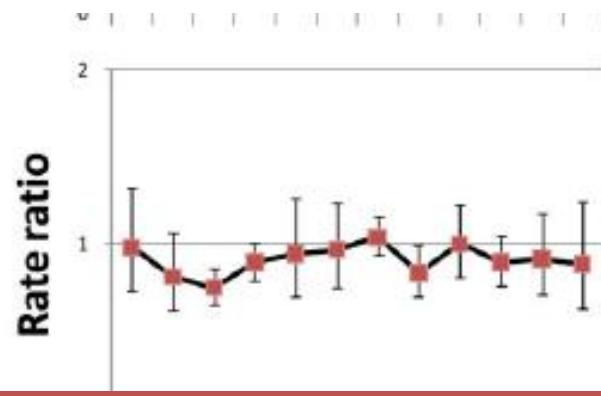
Division of Viral Diseases, National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Atlanta, Georgia

980 • JID 2011:204 (1 October) • BRIEF REPORT

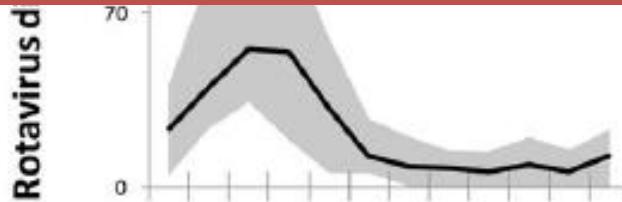
15 to 24 years (C)



(H)



15% de los ingresos evitados y un 20% de los costes evitados se obtuvieron en personas de **5 a 24 años de edad**, es decir, en personas NO vacunadas



Indirect benefits of rotavirus vaccination

« herd immunity or herd protection»

Location	Reduction in rotavirus hospital admissions among	
	Children age-eligible for vaccine	Children not age-eligible for vaccine*
Nationwide, USA ¹	79-81%	69-78%
Nationwide, USA ²	74-85%	41-80%
Queensland, Australia ³	50-70%	30-70%
Nationwide, Belgium ⁴	65-80%	20-64%
Nationwide, Austria ⁵	76-79%	35%
Sao Paulo, Brazil ⁶	56-69%	24%
Nationwide, El Salvador ⁷	79-86%	41-81%

*Typically aged 2-5 years, but age range assessed in each study differed depending on year of vaccine introduction and type assessment.

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2. Yen C et al. Pediatrics. 2011;127:e9–e15.
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4. Raes M et al. Pediatric Infectious Disease Journal. 2011;30: e120–e25.
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6. Safadi MA et al. Pediatric Infectious Disease Journal. 2010;29:1019–1022.
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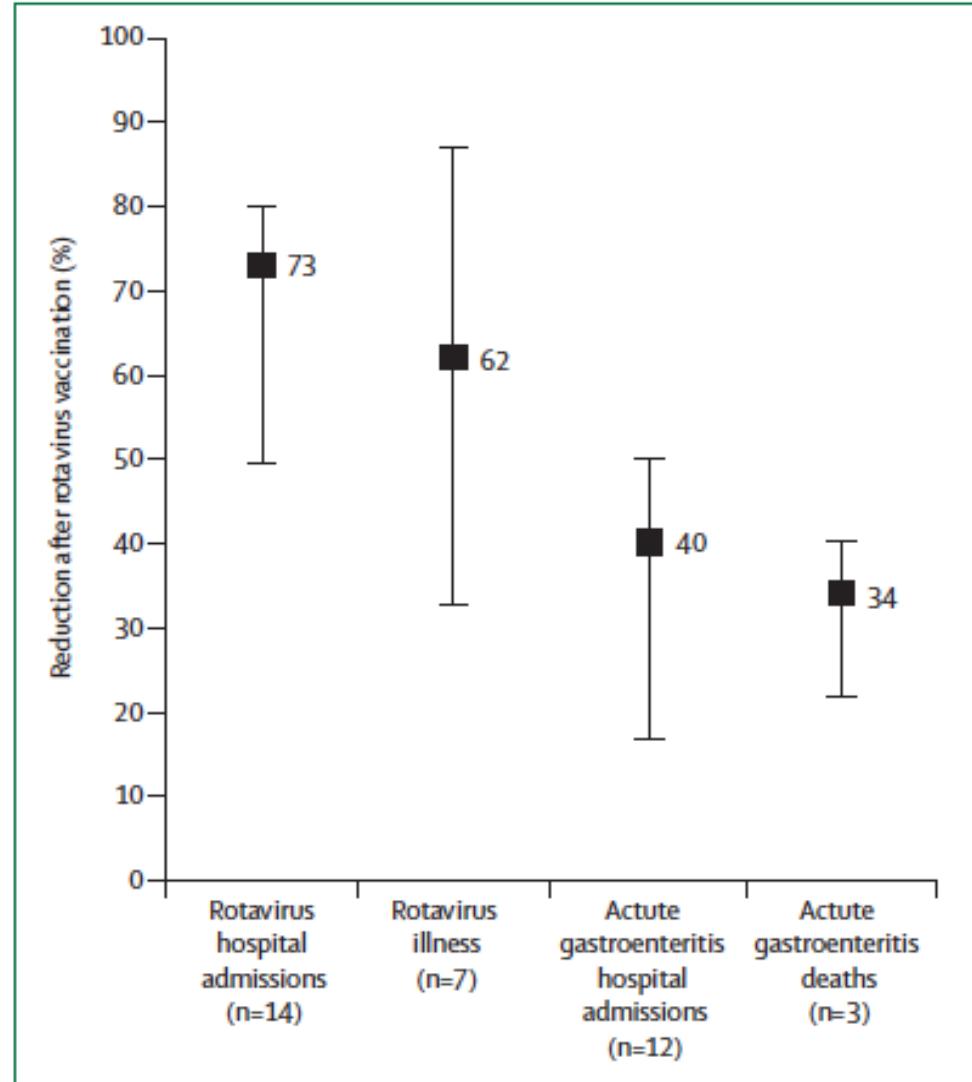
Table derived from Table 2 from: Patel MM, Glass R, Desai R, Tate JE, Parashar UD. Fulfilling the Promise of Rotavirus Vaccines: How Far Have We Come Since Licensure? *The Lancet Infectious Diseases*. 2012;12(7):561-570.

Resumen del valor de la vacunación frente a rotavirus

	Study type	Vaccine	Effectiveness for prevention of RV hospital admissions		Predominant circulating strain
			Full series	Partial series	
High-income					
Houston, TX, USA ⁵⁶	Case-control	RV5	100%	69-81%	G3P[8]
USA (nationwide) ⁵³	Cohort	RV5	100%
New Haven, CT, USA ⁵⁷	Case-control	RV5	94%	93%	G3P[8], NT
Houston, TX, USA ⁵⁸	Case-control	RV5	84%	..	G3P[8]
US military dependants ⁵³	Cohort	RV5	90%	86%	..
Cincinnati, OH, Nashville, TN, and Rochester, NY, USA ⁵⁹	Case-control	RV5	95%	89%	G1-G3, G9, G12
Queensland, Australia ³⁹	Screening	RV5	94%
Central Australia* ⁶⁰	Case-control	RV1	84%	..	G9P[8]
Central Australia*† ⁶¹	Case-control	RV1	51%	..	G2P[4]
Northern Israel‡ ⁶²	Case-control	RV5, RV1	89%
Spain (nationwide)‡ ⁶³	Case-control	RV5, RV1	96%	89%	..
Northwest Brittany, France ⁶⁴	Cohort	RV5	98%
Middle-income					
Northeast Brazil ⁶⁵	Screening	RV1	89-95%	..	G2P[4]
Recife, Brazil ⁶⁵	Case-control	RV1	85%	..	G2P[4]
Belem, Brazil ⁶⁶	Case-control	RV1	40-76%	..	G2P[4]
El Salvador (nationwide) ⁵⁵	Case-control	RV1	76%	51%	G1P[8]
Chiapas, Mexico ⁶⁷	Case-control	RV1	94%	84%	G9P[4]
Low-income					
Four hospitals, Nicaragua ⁵⁴	Case-control	RV5	44%	55%	G2P[4]

RV=rotavirus. RV5=RotaTeq. NT=non-typeable strain. RV1=Rotarix. *Indigenous low-income population in Australia's Northern Territory. †Data are for children younger than 1 year only rather than for all vaccinated children—sample size was small in children older than 1 year, and vaccine effectiveness was lower (9%). ‡Private market only.

Table 3: Postlicensure vaccine effectiveness for prevention of rotavirus hospital admissions



MENSAJES PARA CASA



- **Evidencia científica justifica vacunación frente a rotavirus**
- Las vacunas frente a rotavirus **funcionan**
 - ... **MEJOR** incluso de lo esperado...
 - ... pero sólo funcionan **si se utilizan.**

VIII CONGRESO DE LA SEIP

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