

# Laboratory Detection and Reporting of *Streptococcus agalactiae*

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The presenter states no conflict of interest and has no financial relationship  
to disclose relevant to the content of this presentation.

# OUTLINE

- I. Importance of prenatal screening strategies
- II. Past approaches
- III. Current guidelines
  - A. General indications for prophylaxis
  - B. Laboratory methods and reporting
  - C. Adaptations of molecular approaches
  - D. Antimicrobial susceptibility testing



# Importance of Prenatal Screening



# *Streptococcus agalactiae*

- Colonizes 15-40% of pregnant women

J. Infect. Dis. **143**: 761-766; 1981

Am. J. Obstet. Gynecol. **142**: 617-620; 1982

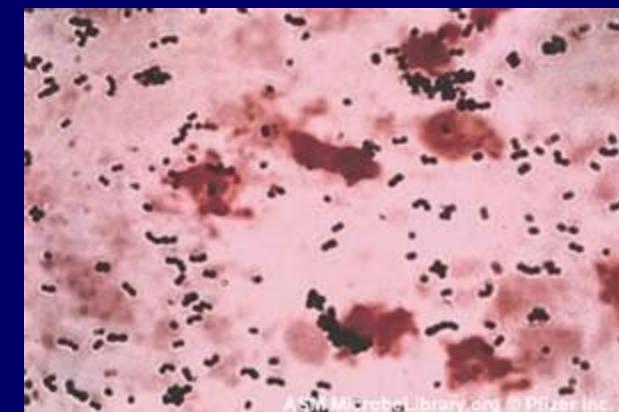
J. Infect. Dis. **145**: 794-799; 1982

J. Infect. Dis. **148**: 802-809; 1983

Obstet. Gynecol. **88**: 811-815; 1996

Obstet. Gynecol. **96**: 498-503; 2000

- Vertical transmission



# *Streptococcus agalactiae*

- Neonatal incidence rate per 1000 live births:

Infection	3.0
Septicemia	2.0
Case fatality	1.0

J. Pediatr. **82:** 707-718; 1973

- Group B streptococcal disease

Early onset	0-72 hours; pneumonia ± bacteremia
Late onset	1-3 months; meningitis

# INTERVENTION

JOURNAL OF CLINICAL MICROBIOLOGY, Mar. 1986, p. 489-492  
0095-1137/86/030489-04\$02.00/0  
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Vol. 23, No. 3

## Reduction of Morbidity and Mortality Rates for Neonatal Group B Streptococcal Disease through Early Diagnosis and Chemoprophylaxis

DANIEL V. LIM,<sup>1\*</sup> WALTER J. MORALES,<sup>2</sup> ANTHONY F. WALSH,<sup>2</sup> AND DENO KAZANIS<sup>3</sup>

*Department of Biology, University of South Florida, Tampa, Florida 33620<sup>1</sup>; Orlando Regional Medical Center, Orlando, Florida 32806<sup>2</sup>; and Orlando Branch Laboratory, Florida Department of Health and Rehabilitative Services, Orlando, Florida 32801<sup>3</sup>*

803 women screened at 36 weeks gestation

173 (21.5%) positive for *S. agalactiae*

80 received intrapartum ampicillin  
93 did not receive antimicrobials

J. Clin. Microbiol. 23: 489-492; 1986

# INTERVENTION

Intrapartum Ampicillin Treatment	Number of Colonized Moms	Number (%) of Colonized Babies
Yes	80	0 (0.0)
No	93	43 (46.2)

# INTERVENTION

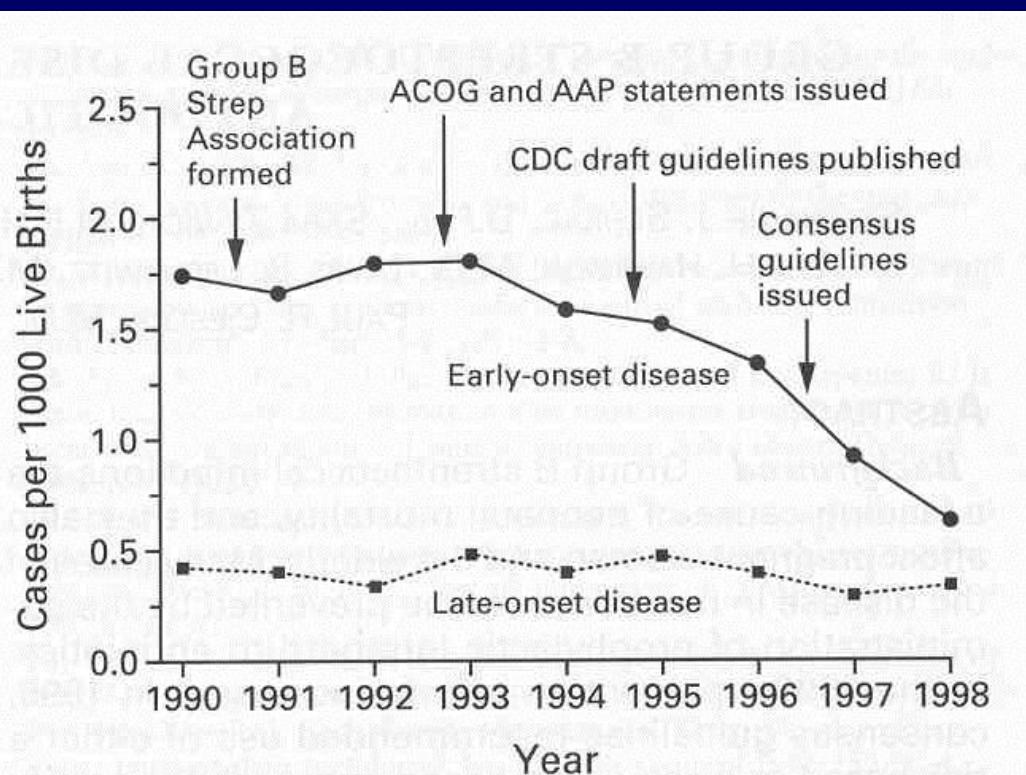
Demographic	Number of Moms	Number of Births	GBS Sepsis/ 1000 Births	
			Incidence	Resultant Fatality
GBS screen + and <b>treated</b> ; GBS screen -	710	710	0.00	0.00
GBS screen + and <b>not treated</b> ; Not screened for GBS	1269	1274	5.49	2.35
<b>Not treated</b> ; Not screened for GBS	3095	3110	2.25	0.32

**Prevention of Perinatal Group B  
Streptococcal Disease: A Public Health  
Perspective**



Second trimester assessment  
Screening- or risk-based

# DISEASE REDUCTION



**Figure 1.** Incidence of Early- and Late-Onset Invasive Group B Streptococcal Disease in Three Active Surveillance Areas (California, Georgia, and Tennessee), 1990 through 1998, and Activities for the Prevention of Group B Streptococcal Disease.

65% reduction in early-onset  
disease prevalence  
from 1993-1998

# The New England Journal of Medicine

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VOLUME 347

JULY 25, 2002

NUMBER 4

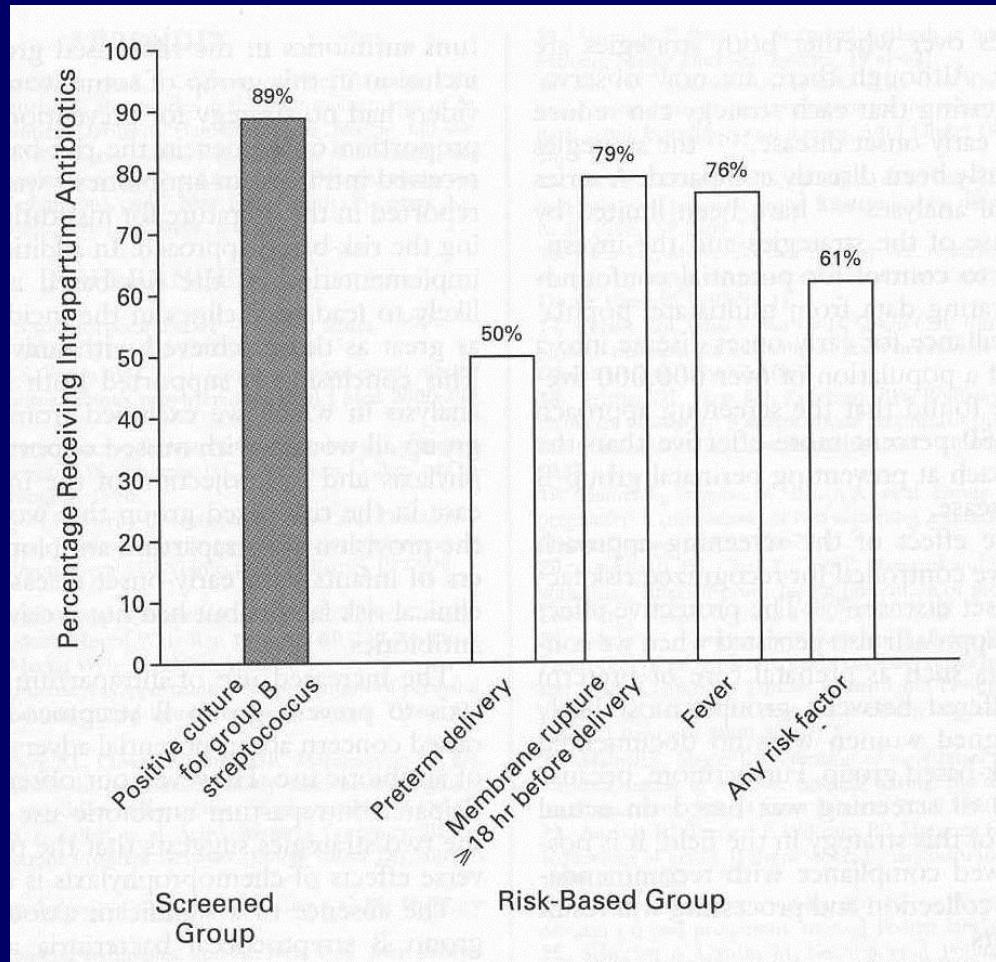


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## A POPULATION-BASED COMPARISON OF STRATEGIES TO PREVENT EARLY-ONSET GROUP B STREPTOCOCCAL DISEASE IN NEONATES

STEPHANIE J. SCHRAG, D.PHIL., ELIZABETH R. ZELL, M.STAT., RUTH LYNFIELD, M.D., AARON ROOME, PH.D.,  
KATHRYN E. ARNOLD, M.D., ALLEN S. CRAIG, M.D., LEE H. HARRISON, M.D., ARTHUR REINGOLD, M.D.,  
KAREN STEFONEK, R.N., M.P.H., GLENDA SMITH, B.S., MELANIE GAMBLE, M.P.H., AND ANNE SCHUCHAT, M.D.,  
FOR THE ACTIVE BACTERIAL CORE SURVEILLANCE TEAM

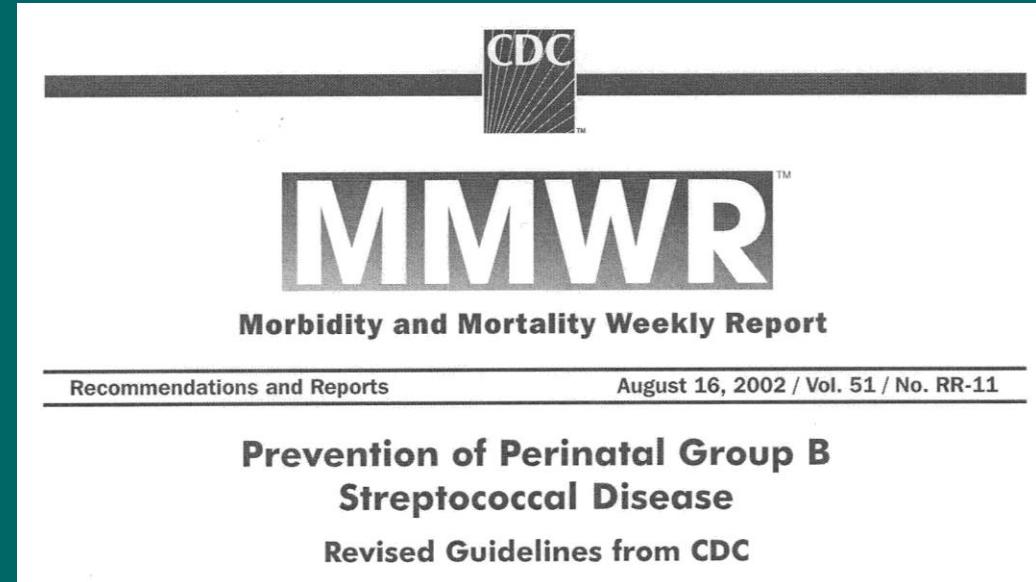
# SCREENING- VERSUS RISK-BASED



Adjusted relative risk for early-onset GBS disease associated with screening approach was 0.48

35- to 37-week assessment

Screening-based



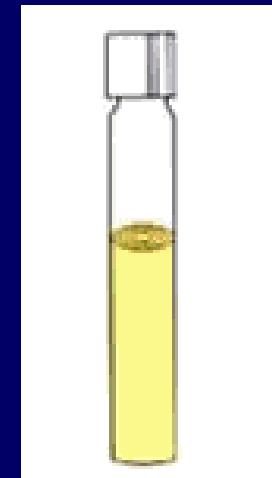
# SCREENING-BASED METHODS



Blood agar  
plate



Blood agar  
plate



+

Todd Hewitt  
(LIM) broth plus  
subculture

*Increases yield 20-35%*

J. Matern. Fet. Med. 7: 172-176; 1998  
Arch. Pathol. Lab. Med. 127: 718-720; 2003

# ADDITIONAL (RECTAL) SAMPLING

Study	Patients	Carriage Rate (%)	Recovery Only by Rectal Sampling (%)
Badri <i>et al.</i> 1977	789	20.5	50.0
Dillon <i>et al.</i> 1982	2540	35.0	51.4
Philipson <i>et al.</i> 1995 <sup>†</sup>	383	20.4	31.1
Platt <i>et al.</i> 1995*	651	16.9	26.4
Quinlan <i>et al.</i> 2000	222	24.3	18.5
Kovavisarach <i>et al.</i> 2007	320	41.9	24.6

J. Infect. Dis. **135**: 308-312; 1977

J. Infect. Dis. **145**: 794-799; 1982

<sup>†</sup>Obstet. Gynecol. **85**: 437-439; 1995

\*Diagn. Microbiol. Infect. Dis. **21**: 65-68; 1995

J. Fam. Pract. **49**: 447-448; 2000

J. Med. Assoc. Thai. **90**: 1710-1714; 2007

# WHO'S SAMPLING??

Investigation	Location	<b>S. agalactiae Culture Sensitivity (%)</b>	
		Patient Collection	Provider Collection
Mercer <i>et al.</i> 1995	Tennessee	91.7 <sup>†</sup>	70.8
Molnar <i>et al.</i> 1997	Ontario	97.4	82.1
Price <i>et al.</i> 2006	Ontario	87.5*	96.9
Arya <i>et al.</i> 2008	Ireland	84.3	94.3

<sup>†</sup> $P < 0.05$

<sup>\*</sup> $P = 0.11$

Am. J. Obstet. Gynecol. **173**: 1325-1328; 1995

Fam. Pract. **14**: 403-406; 1997

J. Obstet. Gynaecol. Can. **28**: 1083-1088; 2006

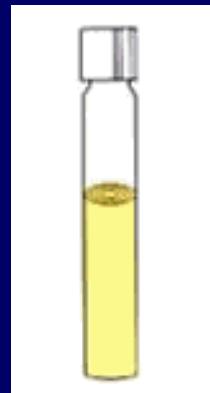
Eur. J. Obstet. Gynecol. Reprod. Biol. **139**: 43-45; 2008

# SCREENING-BASED METHODS



Blood agar  
plate

+



Todd Hewitt  
(LIM) broth plus  
subculture

**87.0% sensitivity**



Blood agar  
plate

+



Carrot broth  
plus subculture

**96.3% sensitivity**

Wheaton Franciscan Laboratory in-house data

# CARROT BROTH (observed at 24h)



Negative for  
*S. agalactiae*

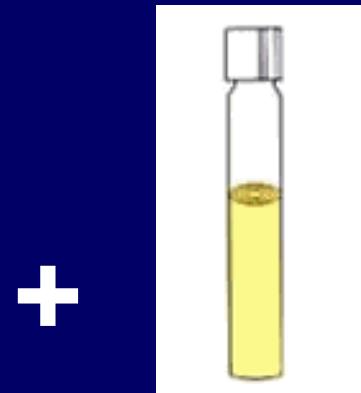


Positive for  
*S. agalactiae*

# SCREENING-BASED METHODS



Blood agar plate



Todd Hewitt  
**(LIM)** broth plus  
subculture

**38.3% resulted on day 1**



Blood agar plate



Carrot broth  
plus subculture

**80.8% resulted on day 1  
( $P < 0.0002$ )**

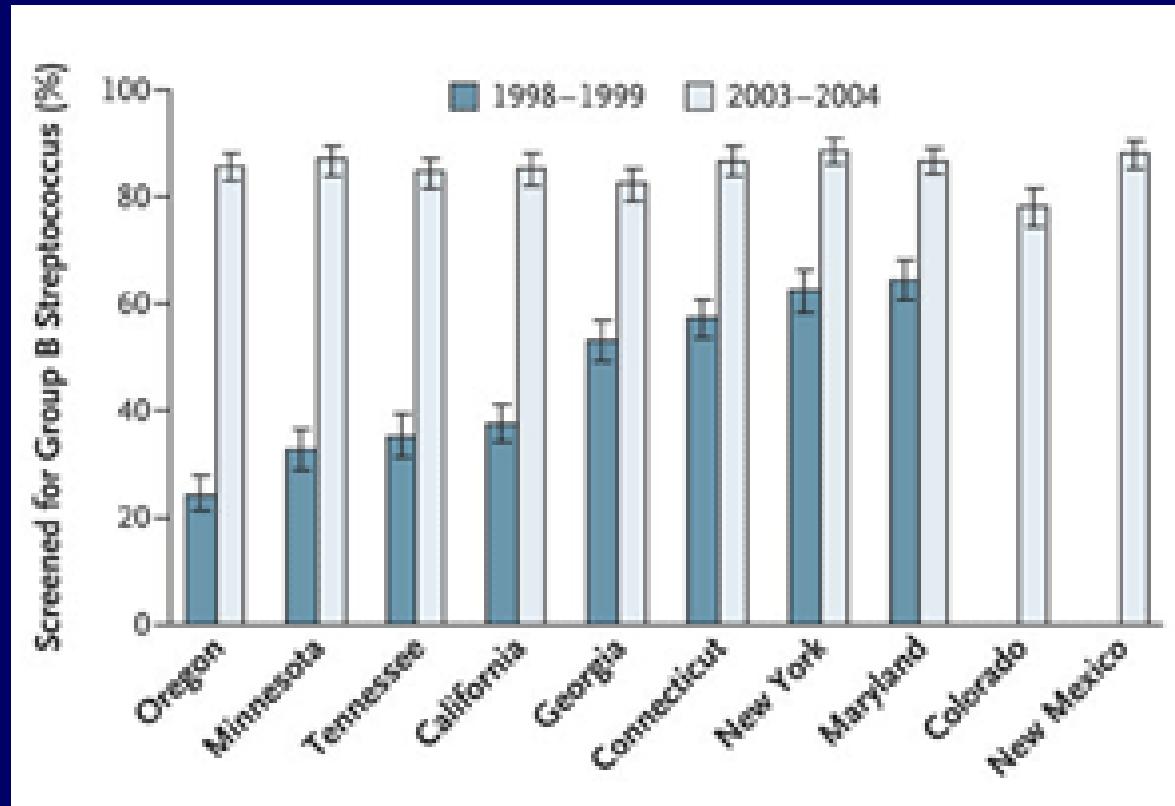
# Is This Working?

ORIGINAL ARTICLE

## Evaluation of Universal Antenatal Screening for Group B Streptococcus

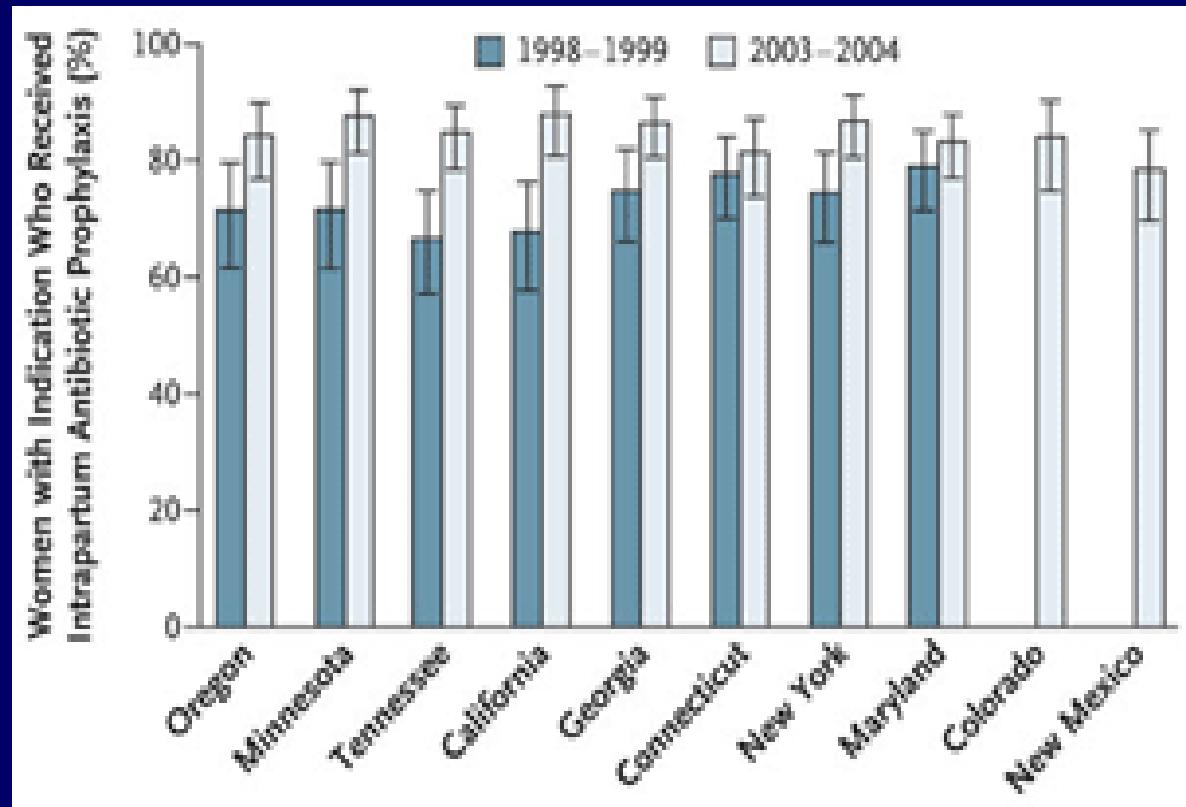
Melissa K. Van Dyke, Ph.D., Christina R. Phares, Ph.D., Ruth Lynfield, M.D.,  
Ann R. Thomas, M.D., Kathryn E. Arnold, M.D., Allen S. Craig, M.D.,  
Janet Mohle-Boetani, M.D., Ken Gershman, M.D., William Schaffner, M.D.,  
Susan Petit, M.P.H., Shelley M. Zansky, Ph.D., Craig A. Morin, M.P.H.,  
Nancy L. Spina, M.P.H., Kathryn Wymore, M.P.H., Lee H. Harrison, M.D.,  
Kathleen A. Shutt, M.S., Joseph Baretta, M.P.H., Sandra N. Bulens, M.P.H.,  
Elizabeth R. Zell, M.Stat., Anne Schuchat, M.D., and Stephanie J. Schrag, D.Phil.

# “SUCCESS” IN SCREENING

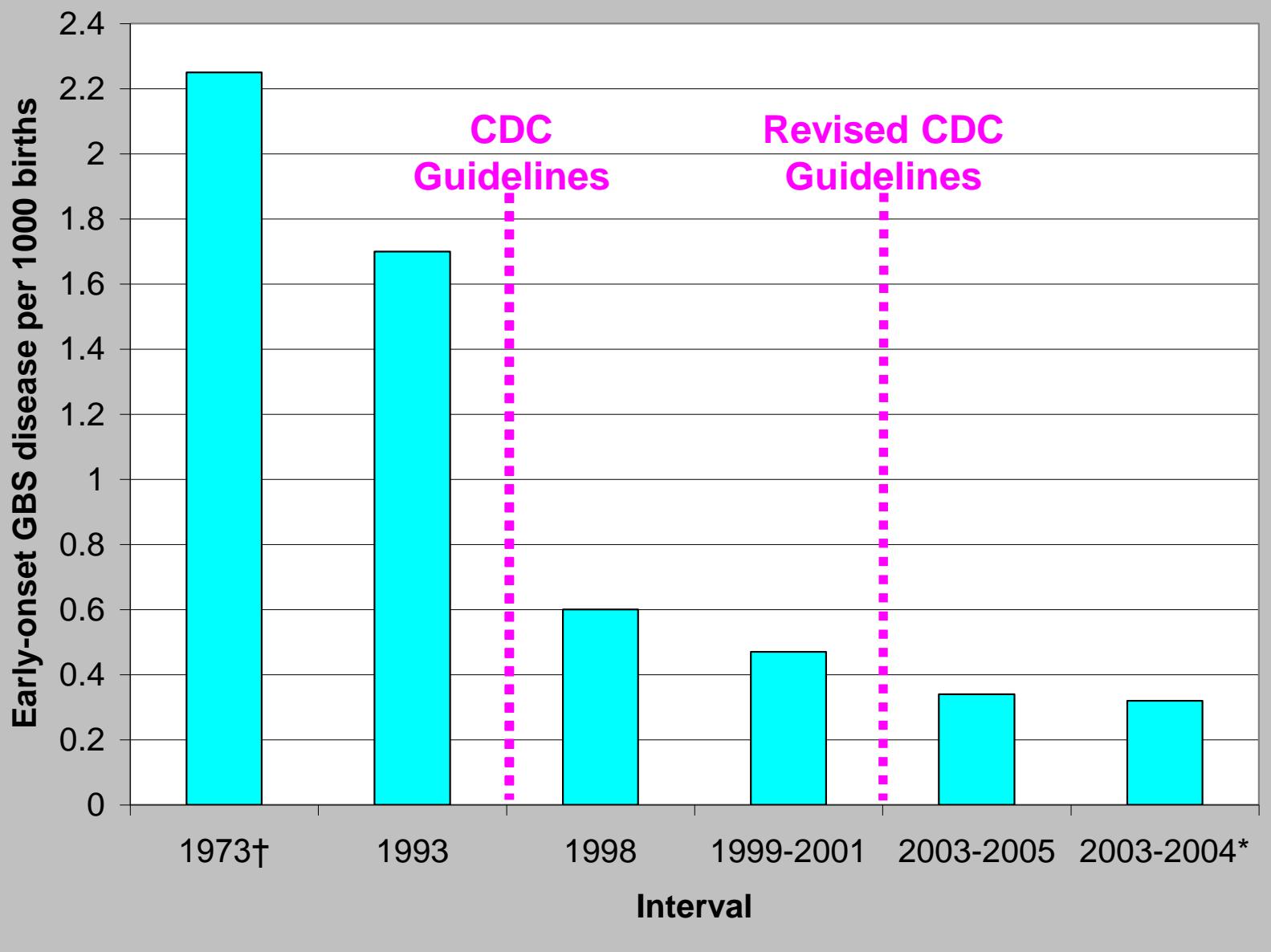


N. Engl. J. Med. 360: 2626-2636; 2009

# “SUCCESS” IN PROPHYLAXIS



N. Engl. J. Med. 360: 2626-2636; 2009



† Estimate  
\* Ten-state surveillance

J. Pediatr. 82: 707-718; 1973  
N. Engl. J. Med. 342: 15-20; 2000

JAMA 299: 2056-2065; 2008  
N. Engl. J. Med. 360: 2626-2636; 2009 24

# DISAPPOINTMENT???

**Table 3.** Implementation of 2002 Recommendations Regarding Intrapartum Chemoprophylaxis, According to Term Status, 2003–2004.\*

Group B Streptococcus Status	Preterm Delivery† (N=962)	Term Delivery (N=6727)
	% (95% CI)	
Positive prenatal screening test before delivery‡		
Total	29.7 (23.9–36.3)	23.9 (22.6–25.2)
Received intrapartum antibiotics		
Overall	84.5 (72.9–91.7)	87.0 (84.9–88.9)
<4 hr between admission and delivery	79.6 (54.8–92.6)	62.7 (56.2–68.8)
≥4 hr between admission and delivery	85.8 (71.7–93.5)	94.0 (92.2–95.5)



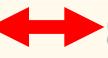
# DISAPPOINTMENT???

**Table 3.** Implementation of 2002 Recommendations Regarding Intrapartum Chemoprophylaxis, According to Term Status, 2003–2004.\*

Group B Streptococcus Status	Preterm Delivery† (N=962)	Term Delivery (N=6727)
	% (95% CI)	
Unknown colonization status§		
Total	54.2 (49.3–59.0)	0.7 (0.5–1.0)
Received intrapartum antibiotics		
Overall	63.4 (57.0–69.4) ←→ 78.5 (63.7–88.4)	
<4 hr between admission and delivery	34.0 (24.3–45.3)	38.9 (8.4–81.5)
≥4 hr between admission and delivery	74.1 (66.7–80.4)	84.3 (69.3–92.7)

# DISAPPOINTMENT???

**Table 3.** Implementation of 2002 Recommendations Regarding Intrapartum Chemoprophylaxis, According to Term Status, 2003–2004.\*

Group B Streptococcus Status	Preterm Delivery† (N=962)	Term Delivery (N=6727)
	% (95% CI)	
History of group B streptococcus bacteriuria or previous infant with group B streptococcus disease		
Total	6.2 (4.3–8.7)	6.7 (6.1–7.5)
Received intrapartum antibiotics		
Overall	73.5 (53.9–86.8) 	80.7 (76.0–84.7)
<4 hr between admission and delivery	59.9 (28.7–84.7)	55.6 (44.5–66.1)
≥4 hr between admission and delivery	74.9 (51.6–89.3)	89.7 (85.0–93.1)

# BIG DISAPPOINTMENT???

Expected 44 to 86 cases of group B streptococcal disease among term infants

→ Observed 116 cases ←

**Table 4.** Characteristics of Mothers Who Delivered at Term and Whose Infants Had Group B Streptococcal Disease, 2003–2004.

Characteristic	Mothers Who Delivered at Term and Whose Infants Had Group B Streptococcal Disease (N=189)
	no. (%)
Screened	155 (82.0)
Positive for group B streptococcus	37 (19.6)
Negative for group B streptococcus	116 (61.4) 
Unknown colonization status	2 (1.1)

# Need improved diagnostics

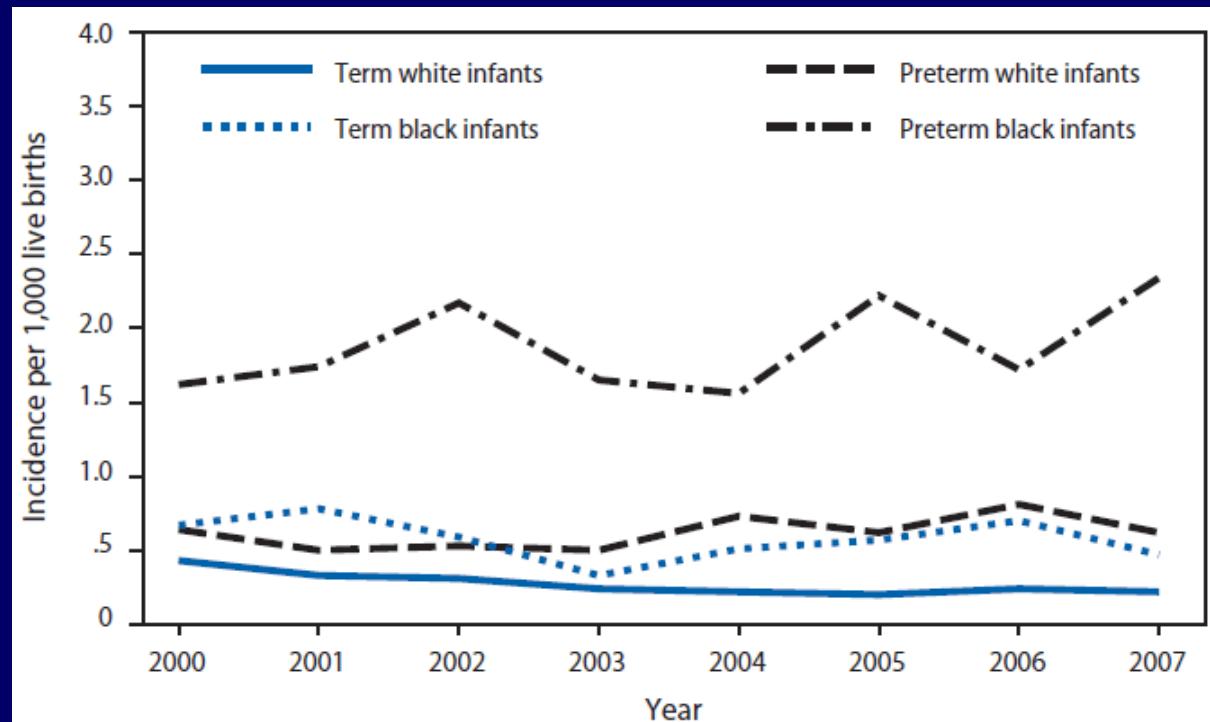
At same time, demographics may benefit from rapid & accurate diagnostics

# BENEFIT FROM A RAPID RESULT

- Increased attack rates and mortality in low birth weight neonates

**TABLE 1.** NUMBER OF CASES OF EARLY-ONSET NEONATAL INVASIVE GROUP B STREPTOCOCCAL DISEASE AND CASE FATALITY RATES ACCORDING TO GESTATIONAL AGE IN SELECTED COUNTIES IN THE UNITED STATES, 1993 TO 1998.

GESTATIONAL AGE	No. (% OF EARLY-ONSET CASES)	CASE FATALITY RATE (%)*
≤33 wk	137 (9)	30
34–36 wk	116 (7)	10
≥37 wk	1247 (83)	2



N. Engl. J. Med. 342:  
15-20; 2000

MMWR. 59 (RR-10):  
1-32; 2010

# BENEFIT FROM A RAPID RESULT

- Increased attack rates and mortality in low birth weight neonates
- Inadequate/no prenatal care

Higher probability in African Americans

Increased disease in those with inadequate care

Increased disease in African American neonates

Obstet. Gynecol. **87**: 575-580; 1996

Obstet. Gynecol. **89**: 28-32; 1997

# BENEFIT FROM A RAPID RESULT

- Increased attack rates and mortality in low birth weight neonates
- Inadequate/no prenatal care
- Moms who screen negative at 35-37 weeks, but are colonized at parturition (estimated 4-9%)

Pediatrics 115: 1240-1246; 2005  
J. Infect. Dis. 148: 802-809; 2005

# COMMERCIAL PCR

- Rapid detection of *S. agalactiae* DNA in vaginal/rectal specimens from prepartum or intrapartum women (direct swab)
- 86-94% sensitivity (**LIM** broth reference)

Clin. Infect. Dis. **39**: 1129-1135; 2004

# PERFORMANCE INDICES

Carrot broth-enhanced PCR 33.0% detection

LIM broth-enhanced PCR 30.5% detection

Carrot broth culture 29.6% detection

Parameter	Carrot Broth PCR	LIM Broth PCR
Sensitivity (%)	100.0	92.5
Negative predictive value (%)	100.0	96.4
Unresolved rate (%)	0.0	0.5
Processing time/specimen (min)	5.1	5.1

# PCR-POSITIVE SPECIMENS; n = 67

Parameter	Timepoint of Carrot Broth Culture Observation	
	Overnight Incubation	Final Subculture Result
Positive culture	34	60
Sensitivity (%)	50.7	89.6
Negative predictive value (%)	80.5	95.1

# COMMERCIAL PCR

- Rapid detection of *S. agalactiae* DNA in vaginal/rectal specimens from prepartum or intrapartum women (direct swab)
- 86-94% sensitivity (**LIM** broth reference)

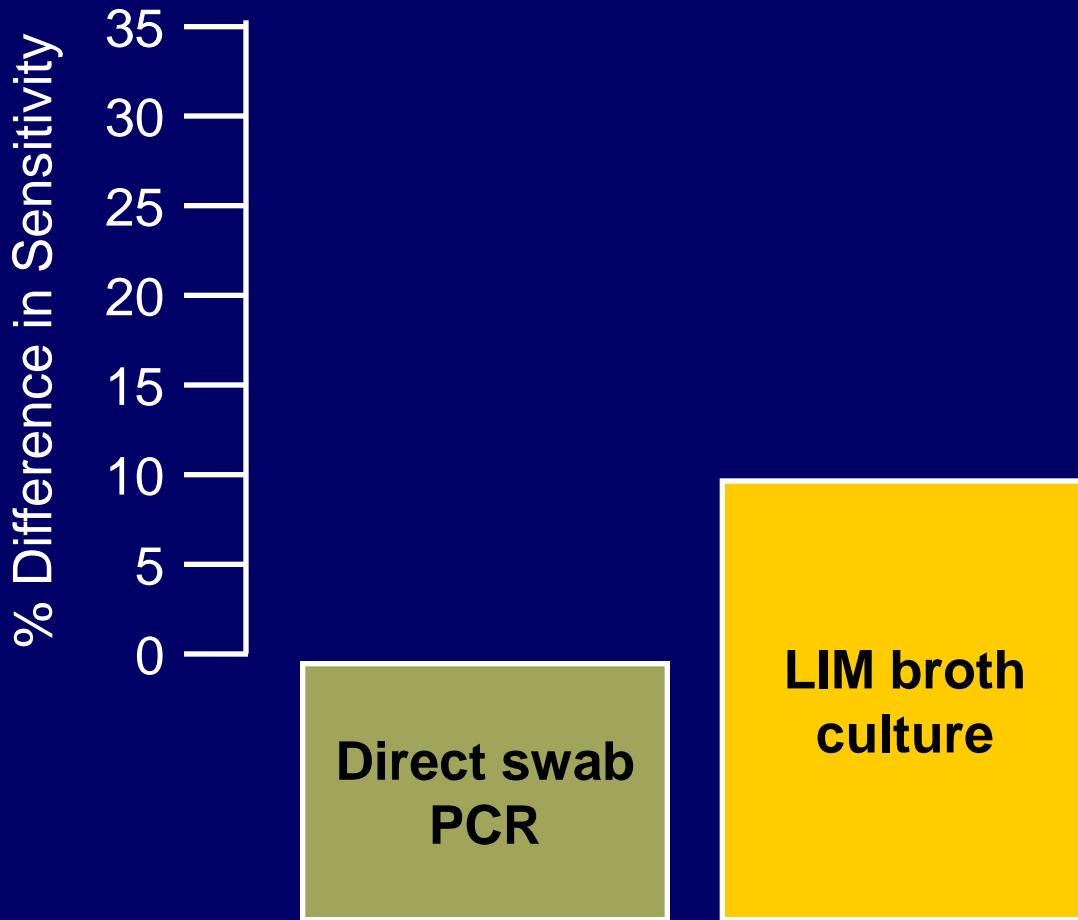
Clin. Infect. Dis. **39**: 1129-1135; 2004

- 56-59% sensitivity (**carrot** broth reference)

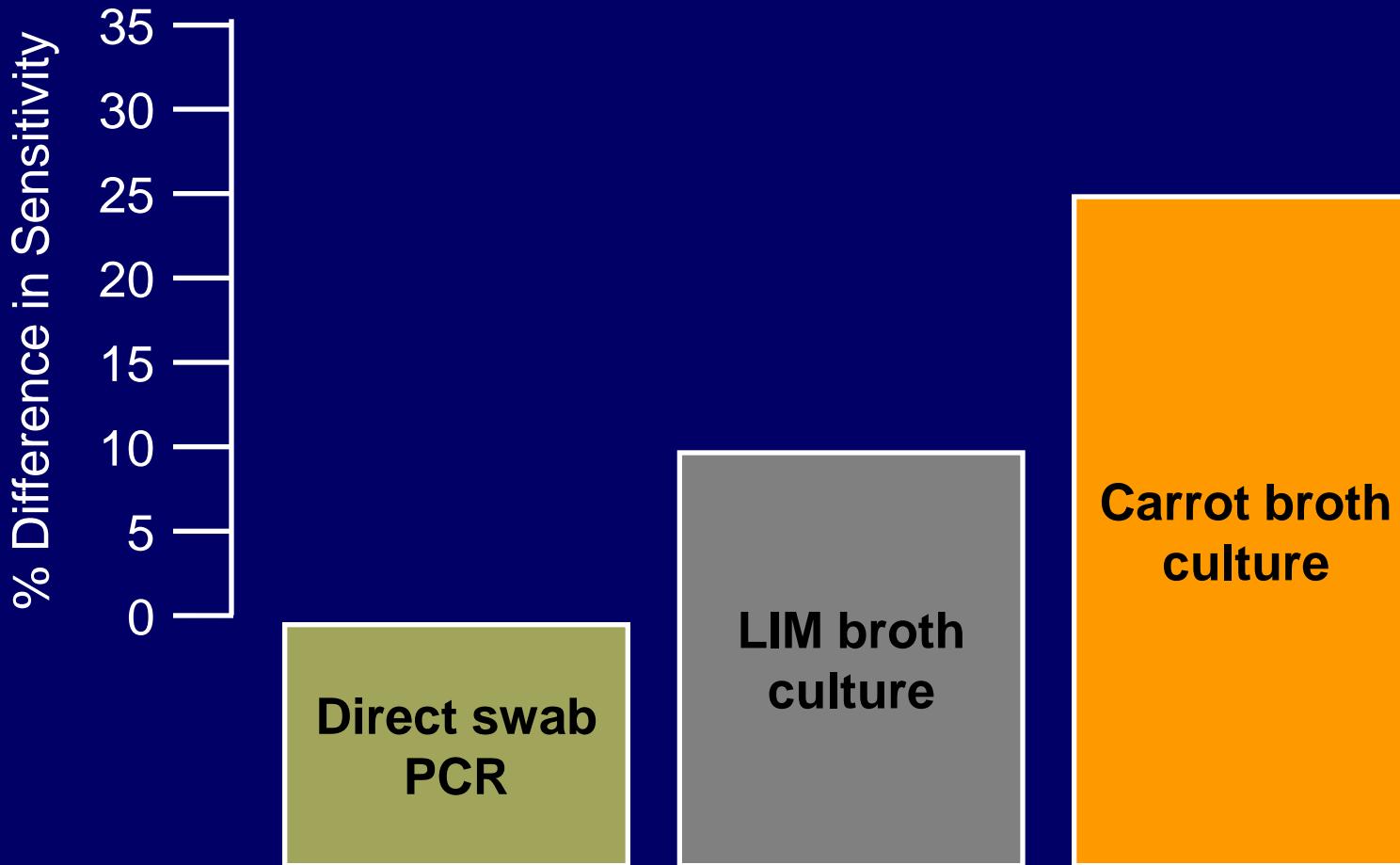
J. Clin. Microbiol. **46**: 3615-3620; 2008

J. Clin. Microbiol. **48**: 4495-4500; 2010

# WHY??

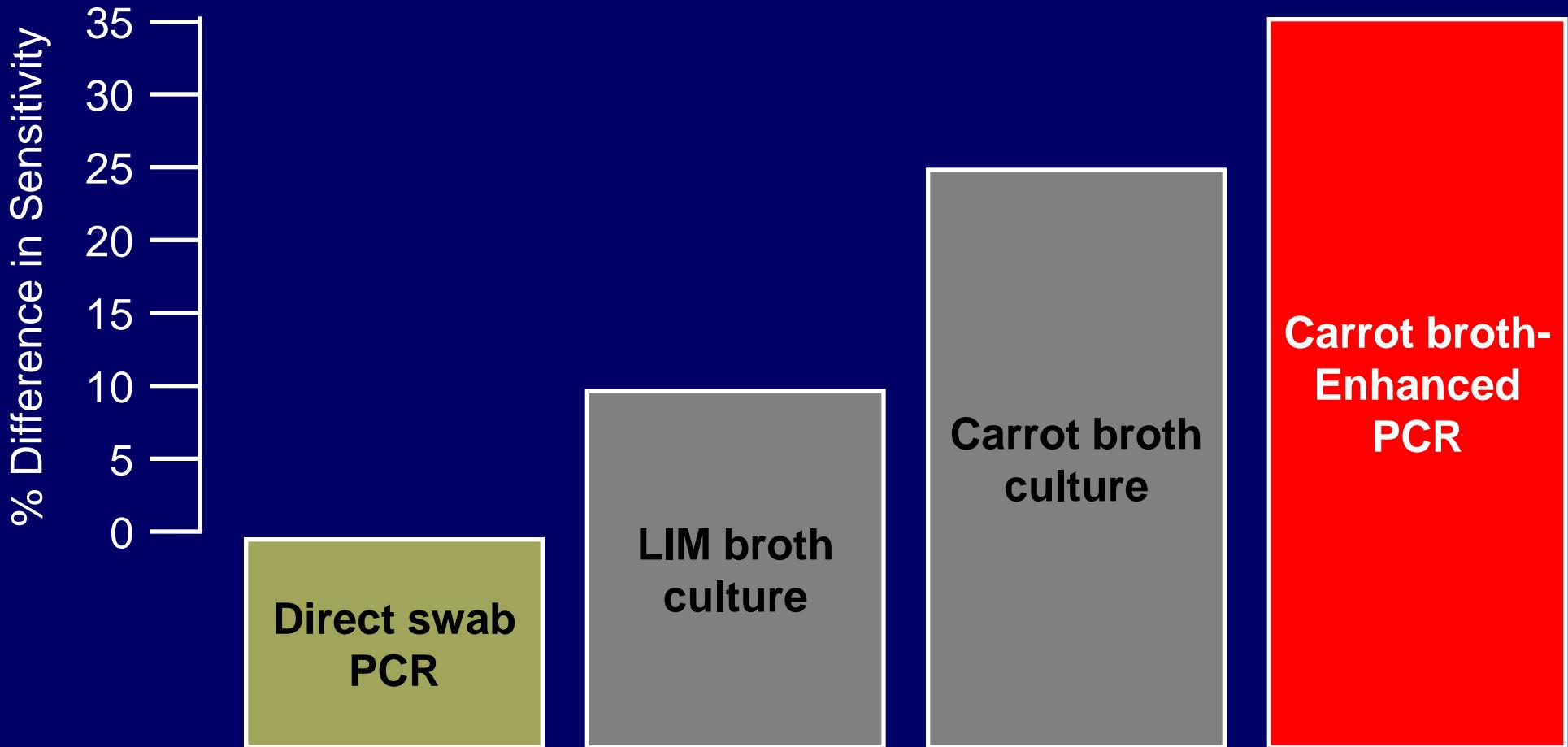


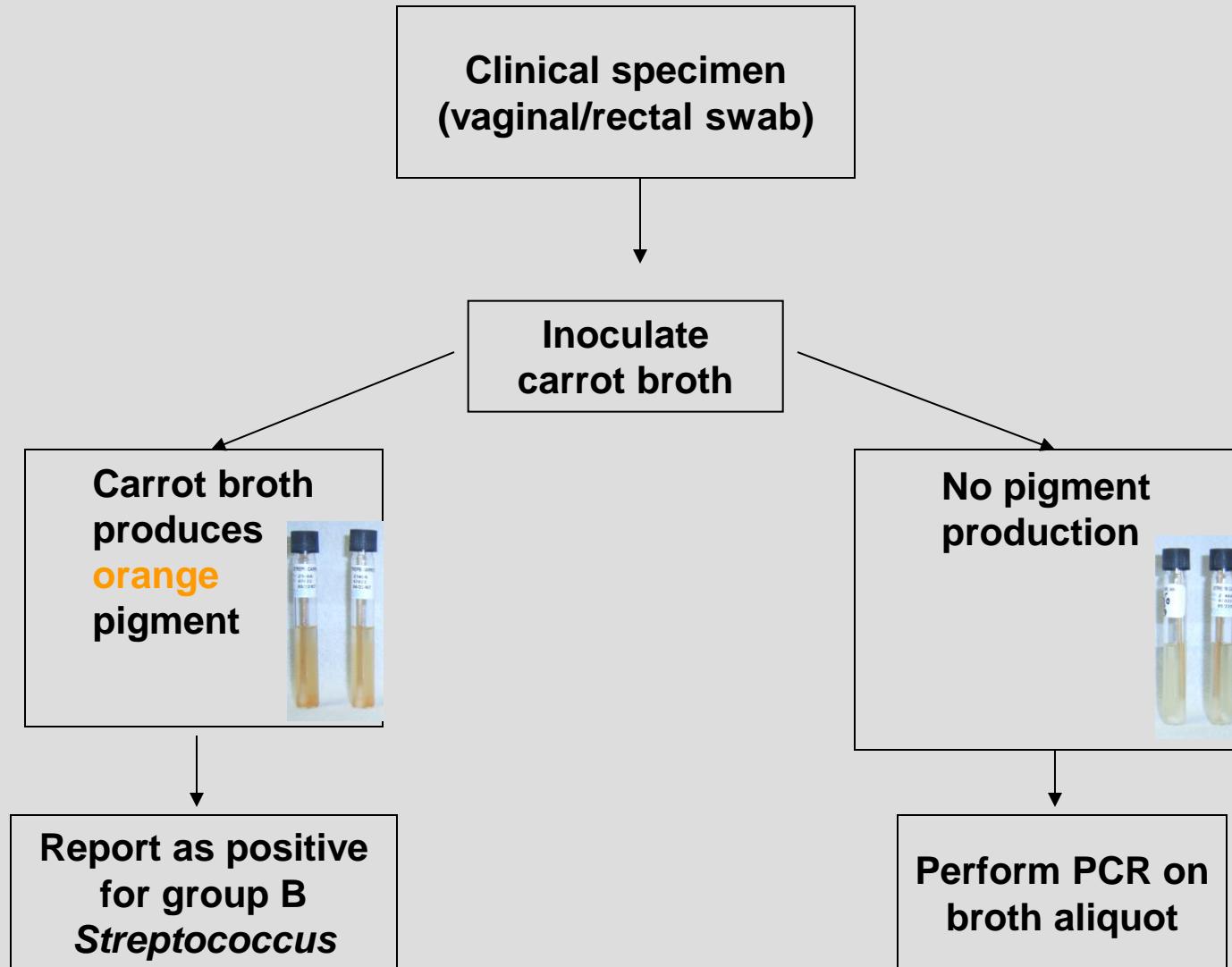
# WHY??



Wheaton Franciscan Laboratory in-house data

# WHY??





**CAN THIS BECOME  
MORE RAPID??**

# *IN VITRO* EXPERIMENTATION

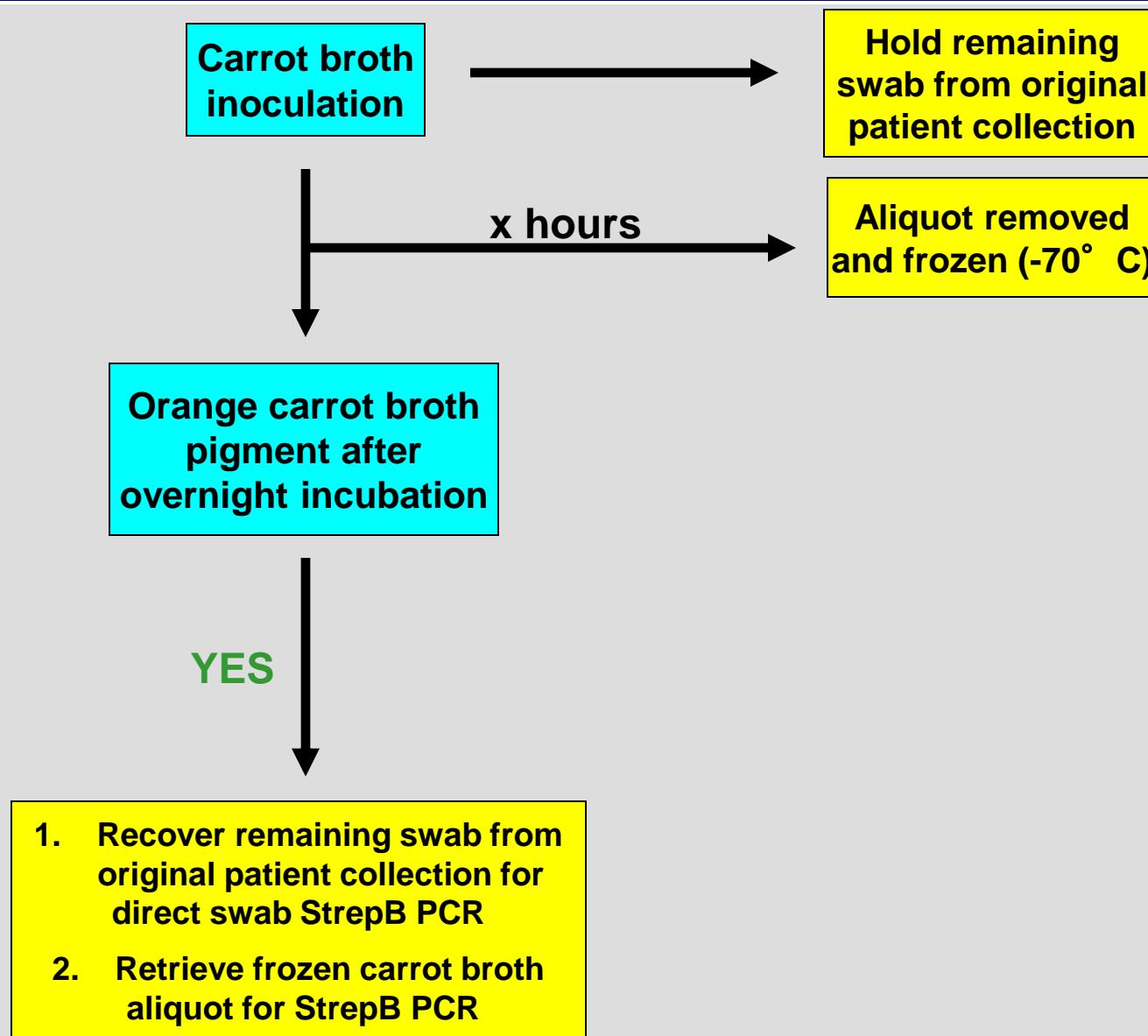
- Inoculate carrot broth tubes with  $10^3$ ,  $10^2$ ,  $10^1$  *S. agalactiae*
- Mock inoculation with  $10^9$  flora; simulating...
  - Anaerobic flora
  - Gastrointestinal flora
  - Urogenital flora
  - Pathogenic flora
- Collect 500- $\mu$ L aliquots at specified intervals for carrot broth-enhanced PCR

# CARROT BROTH-ENHANCED PCR

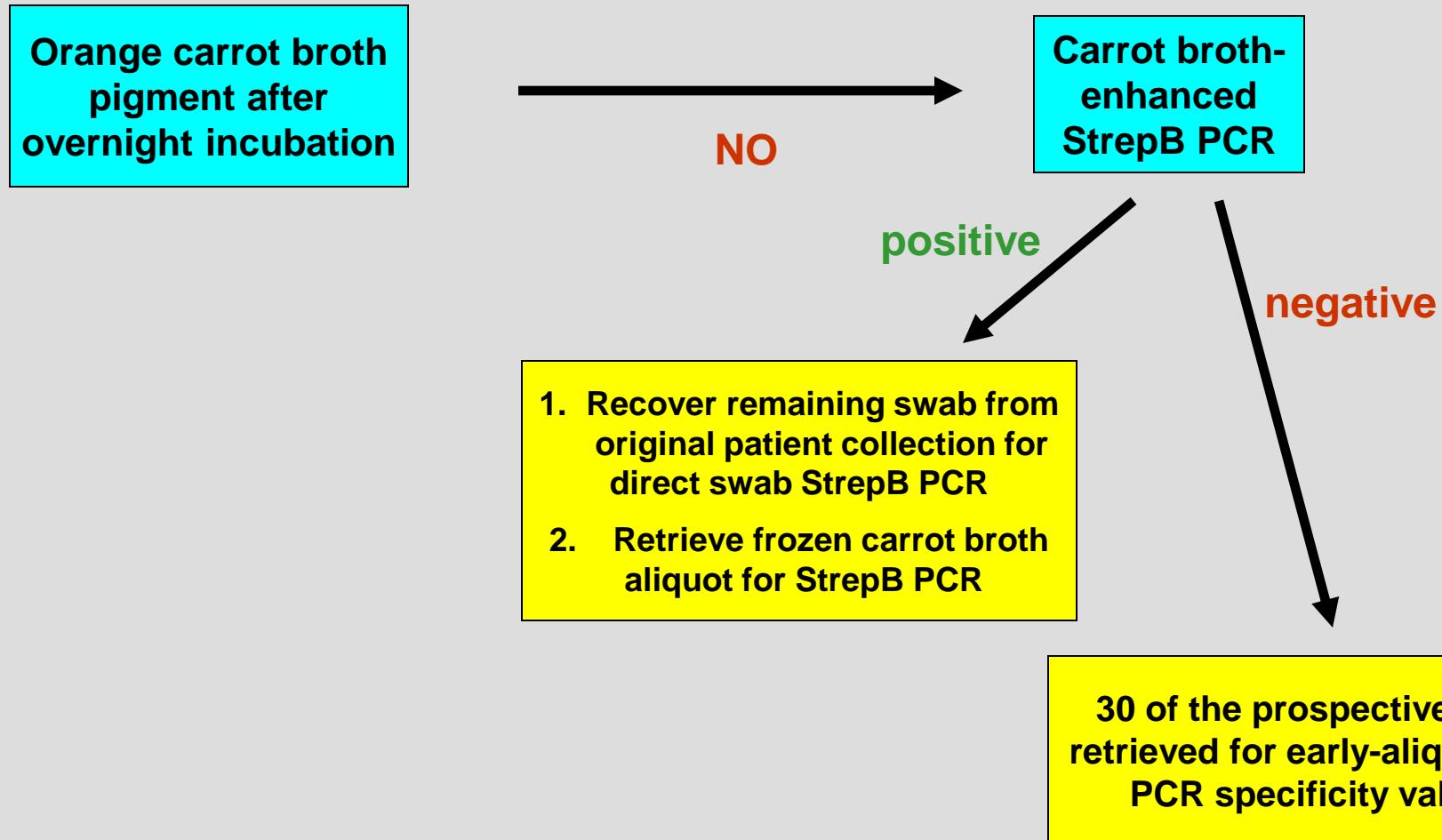
<i>S. agalactiae</i> Inoculum	Percentage Positive				
	<i>Time of aliquot collection (hours)</i>				
	2	4	6	12	24
10 <sup>1</sup>	0.0	33.3	41.7	25.0	33.3
10 <sup>2</sup>	58.3	66.7	91.7	ND	ND
10 <sup>3</sup>	100.0	100.0	100.0	ND	ND

ND; not determined

# CLINICAL EXPERIMENTATION



# CLINICAL EXPERIMENTATION



# CLINICAL EXPERIMENTATION

Number of Specimens	Early-aliquot Carrot Broth-enhanced PCR		% Positive from Remnant Direct Swab PCR	P value
	Collection Interval (h)	% Positive		
33	< 3.00	54.5	66.7	0.31
35	3.00-3.99	40.0	54.3	0.23
35	4.00-4.99	51.4	48.6	0.81
41	5.00-5.99	73.2	65.9	0.47
39	6.00-6.99	82.1	46.2	0.0009
44	> 7.00	86.3	56.8	0.002
Total (227)		66.1	56.4	0.03

# POSITIVE CARROT BROTH CULTURE

Number of Specimens	Early-aliquot Carrot Broth-enhanced PCR		% Positive from Remnant Direct Swab PCR	<i>P</i> value
	Collection Interval (h)	% Positive		
12	< 3.00	83.3	91.7	ND
12	3.00-3.99	50.0	75.0	ND
10	4.00-4.99	80.0	80.0	ND
19	5.00-5.99	94.7	89.5	ND
13	6.00-6.99	100.0	69.2	ND
10	> 7.00	100.0	70.0	ND
Total (76)		85.5	80.3	0.39

# NEGATIVE CARROT BROTH CULTURE

Number of Specimens	Early-aliquot Carrot Broth-enhanced PCR		% Positive from Remnant Direct Swab PCR	<i>P</i> value
	Collection Interval (h)	% Positive		
21	< 3.00	38.1	52.4	0.35
23	3.00-3.99	34.7	43.5	0.55
25	4.00-4.99	40.0	36.0	0.77
22	5.00-5.99	54.5	45.5	0.55
26	6.00-6.99	73.1	34.6	0.005
34	> 7.00	82.4	52.9	0.01
Total (151)		56.2	44.4	0.04



# MMWR<sup>TM</sup>

**Morbidity and Mortality Weekly Report**

[www.cdc.gov/mmwr](http://www.cdc.gov/mmwr)

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Recommendations and Reports

November 19, 2010 / Vol. 59 / No. RR-10

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## **Prevention of Perinatal Group B Streptococcal Disease**

**Revised Guidelines from CDC, 2010**

35- to 37-week assessment

Screening-based

# INDICATIONS FOR PROPHYLAXIS

- Previous infant with invasive early-onset disease
- *S. agalactiae* bacteriuria during any trimester of current pregnancy
- Positive *S. agalactiae* vaginal/rectal screening culture in late gestation during current pregnancy
- Unknown *S. agalactiae* status at labor **PLUS** one:

Delivery at < 37 weeks' gestation

Amniotic membrane rupture  $\geq$  18 hours

Intrapartum temperature  $\geq$  100.4° F

Positive intrapartum nucleic acid amplification test

# SPECIMEN COLLECTION/TRANSIT

- Lower vaginal, then rectal collection
  - 35-37 weeks' gestation; can be self-collected
  - Cervical, perianal, perirectal not acceptable
- Swabs placed into non-nutritive transport medium
  - Recovery decreases over 1-4 days (room temp)
  - Refrigerate swabs, if feasible
- Clinicians indicate if patient possesses allergy to penicillin or cephem agent

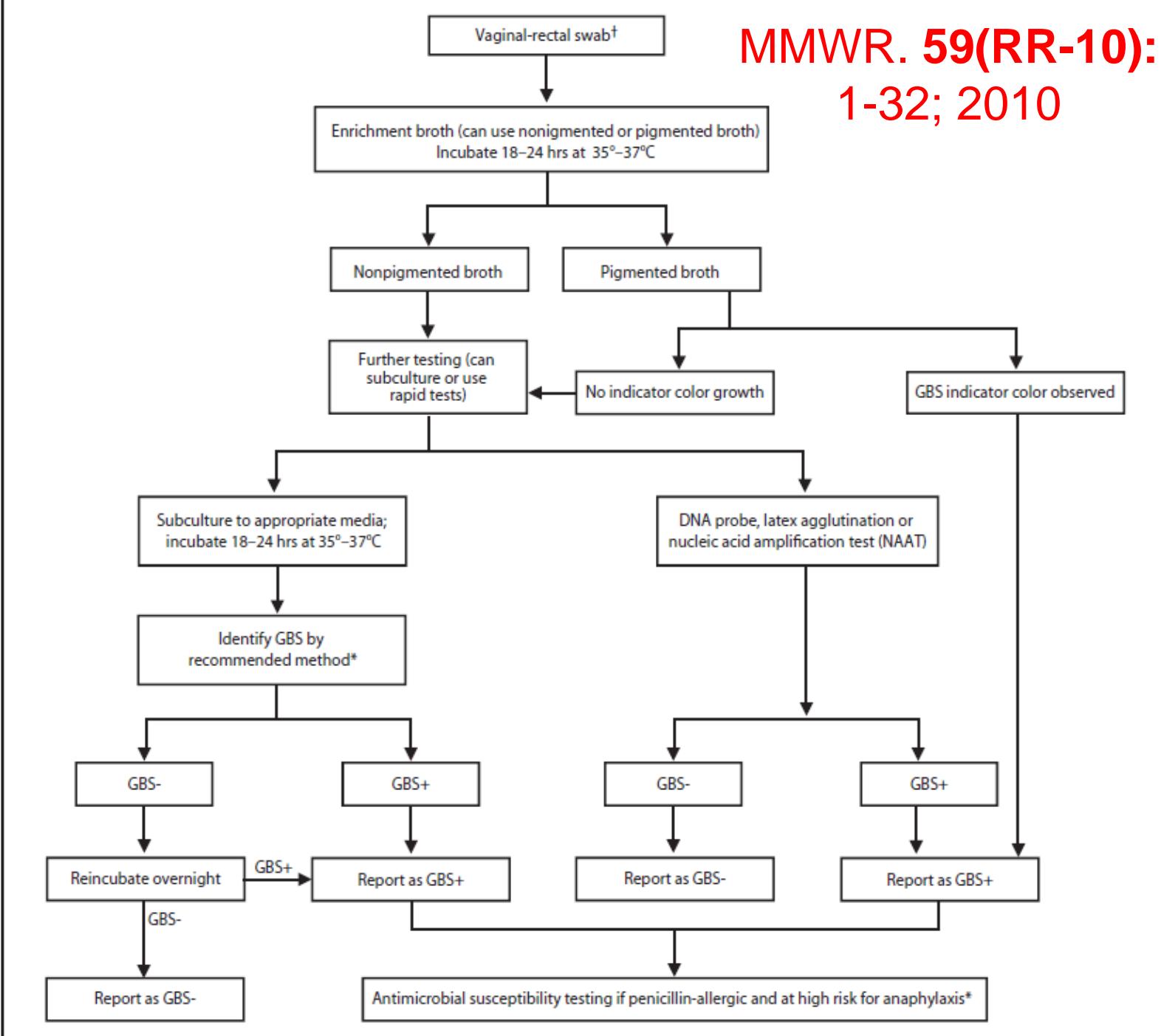
# SPECIMEN PROCESSING

- Selective broth medium (Todd-Hewitt base)
    - LIM broth
    - Transvag broth
  - Alternative selective media can be chromogenic
    - Carrot broth
    - Granada biphasic broth
  - 18-24 hours in 35-37C ambient air or 5% CO<sub>2</sub>
  - Direct plating may be included
    - Lower sensitivity than broth enrichment
    - Should not be used as sole means of recovery
- MMWR. 59 (RR-10): 1-32; 2010

# RESULTS AND INTERPRETATION

- Selective broths subcultured to appropriate agar(s)
- Non-pigmented chromogenic broths subcultured to appropriate agar(s)
- Positive identification may be derived from:
  - Biochemical or probe testing of isolated growth
  - Pigmented broth ( $\beta$ -hemolytic *S. agalactiae*)
  - Probe testing of selective broth
  - Nucleic acid amplification testing of selective broth
  - Latex agglutination of selective broth

**MMWR. 59(RR-10):  
1-32; 2010**



# DIRECT MOLECULAR DETECTION?

"Accurate results are more important than rapid turnaround time for antenatal screening."

MMWR. 59 (RR-10): 1-32; 2010

College of American Pathologists MIC.64817

"A pre-enrichment step using a selective broth enrichment culture is performed for antepartum (35-37 weeks gestation) vaginal/rectal swab screening for Group B streptococci (GBS) colonization by nucleic acid amplification testing (NAAT)."

# ANTIMICROBIAL SUSCEPTIBILITY

- Disk diffusion or broth microdilution performed on antenatal *S. agalactiae* isolates from women at risk for anaphylaxis (related to penicillin or cephem)

Anaphylaxis

Angiodedema

Respiratory distress

Urticaria

- Inducible clindamycin testing on erythromycin-resistant *S. agalactiae*
- CLSI M100 document recommends suppression of erythromycin susceptibility testing data

# INDUCIBLE CLINDAMYCIN RESISTANCE

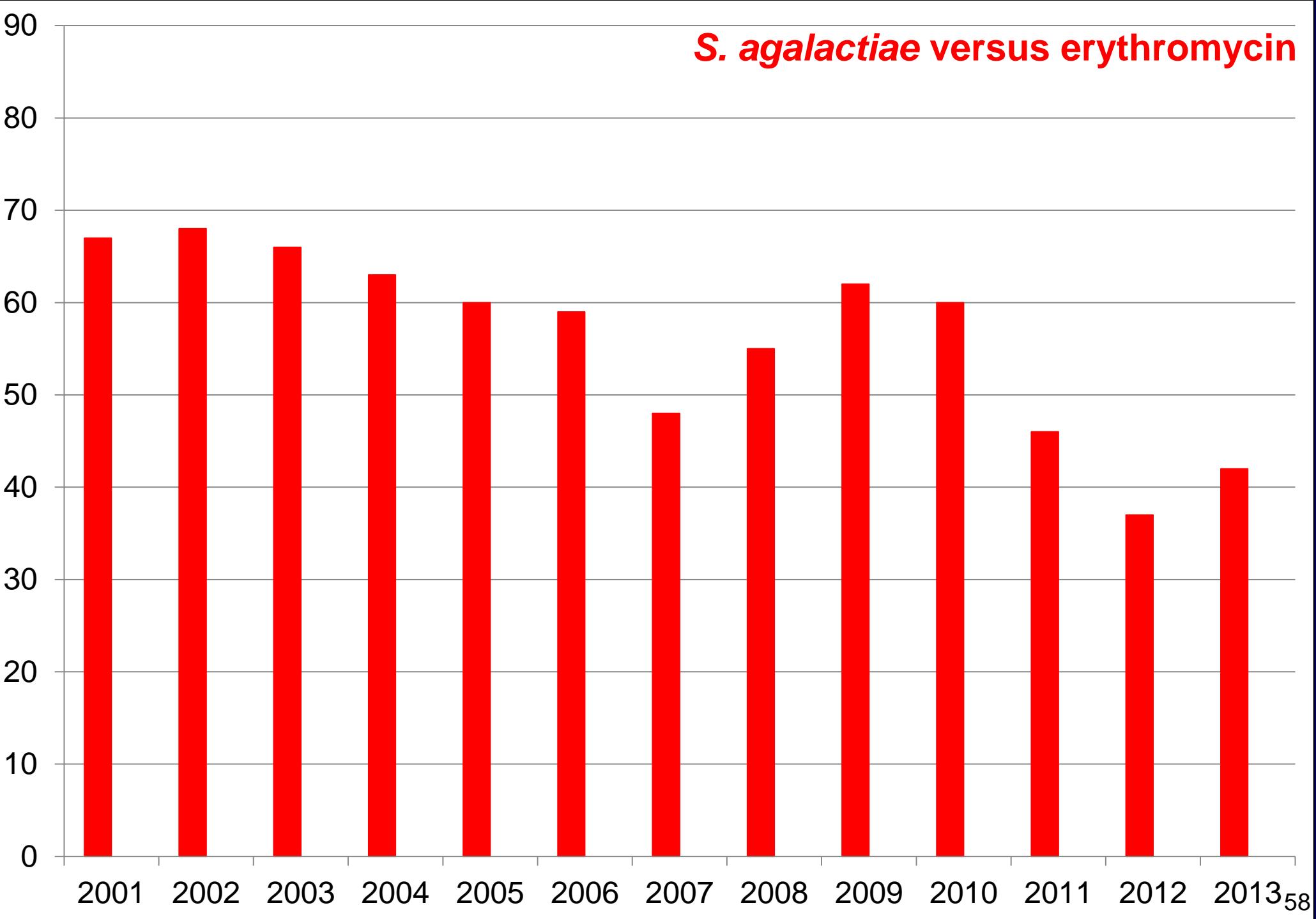
- CDC recommends “D-test” on erythromycin-R/clindamycin-S isolates of *S. agalactiae*; allows for performance on other validated AST systems
- 2 µg clindamycin disk  
15 µg erythromycin disk  
12 millimeters apart
- Mueller-Hinton w/blood  
35C; 5% CO<sub>2</sub>  
20-24 hours



MMWR. 59 (RR-10): 1-32; 2010

*S. agalactiae* versus erythromycin

Percentage Susceptible

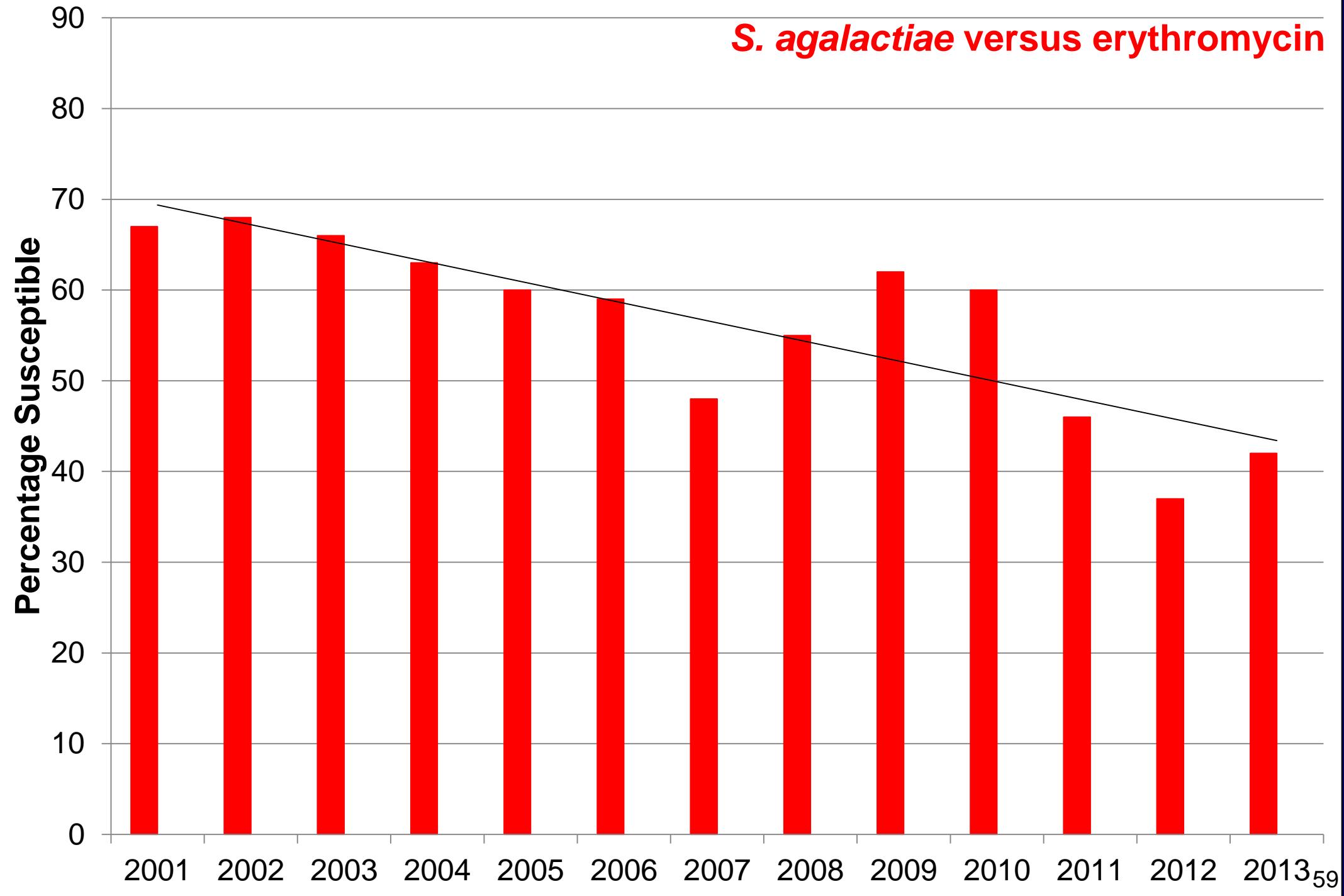


*S. agalactiae* versus erythromycin

Percentage Susceptible

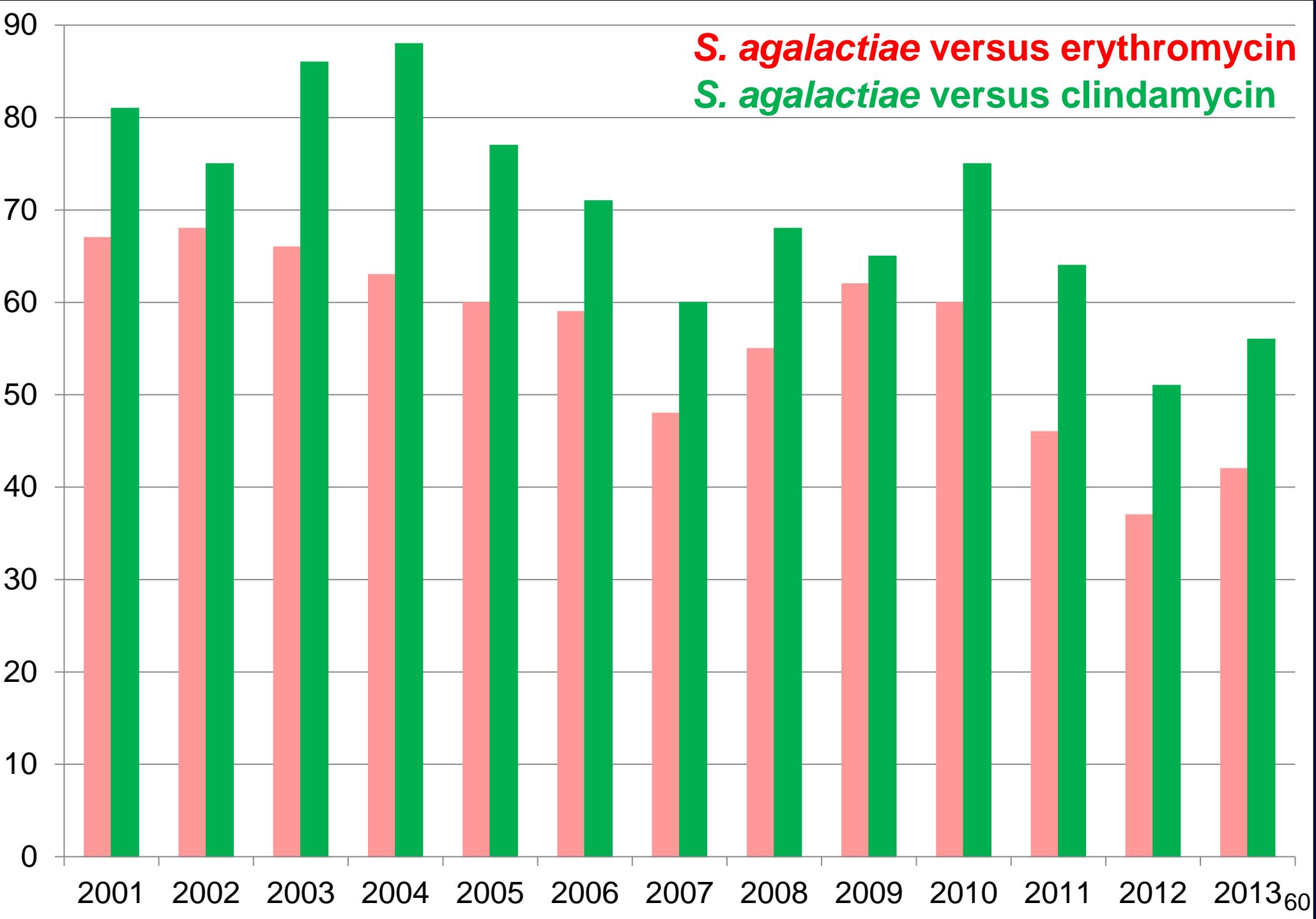
90  
80  
70  
60  
50  
40  
30  
20  
10  
0

2001 2002 2003 2004 2005 2006 2007 2008 2009 2010 2011 2012 2013<sub>59</sub>



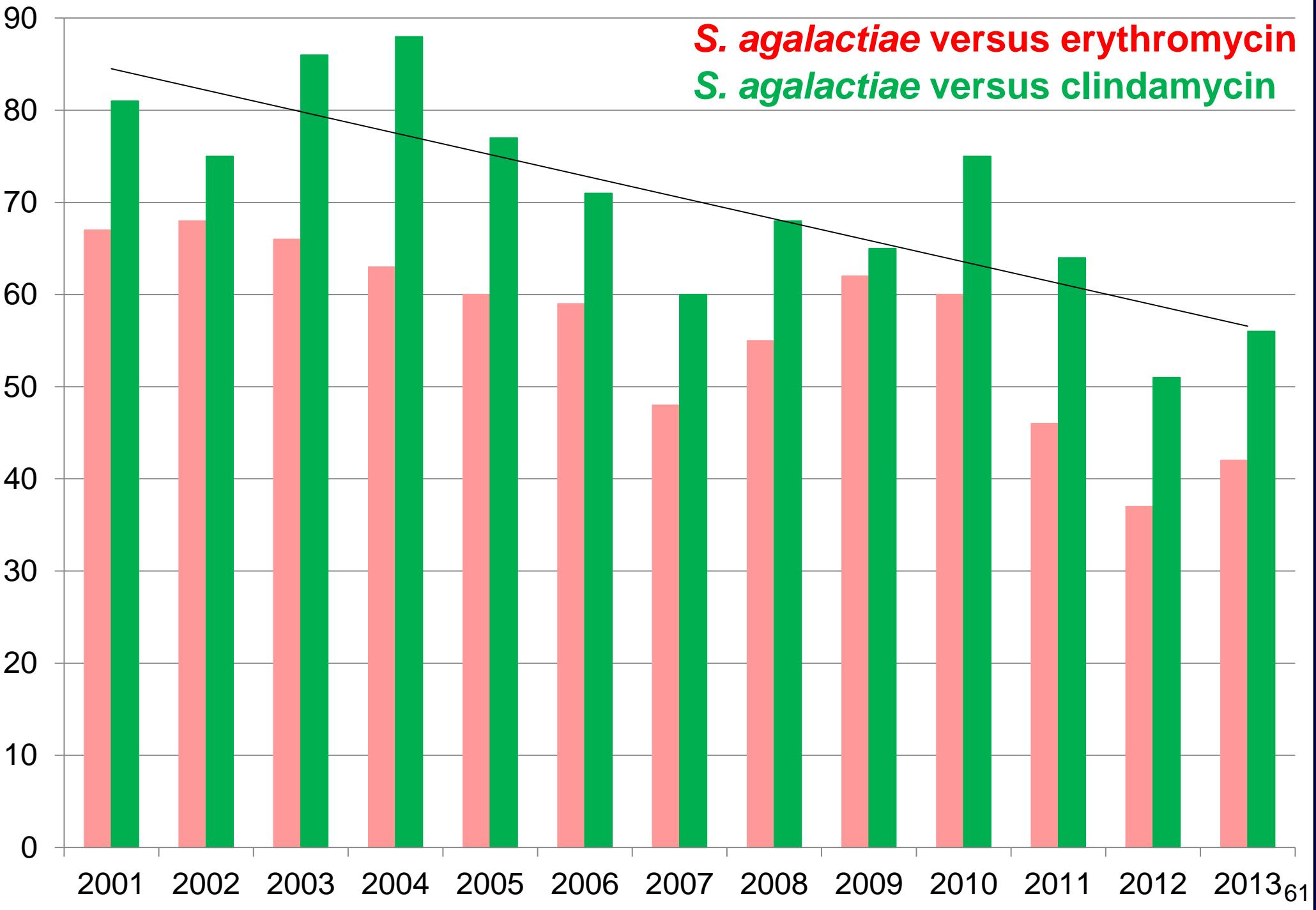
***S. agalactiae* versus erythromycin**  
***S. agalactiae* versus clindamycin**

**Percentage Susceptible**

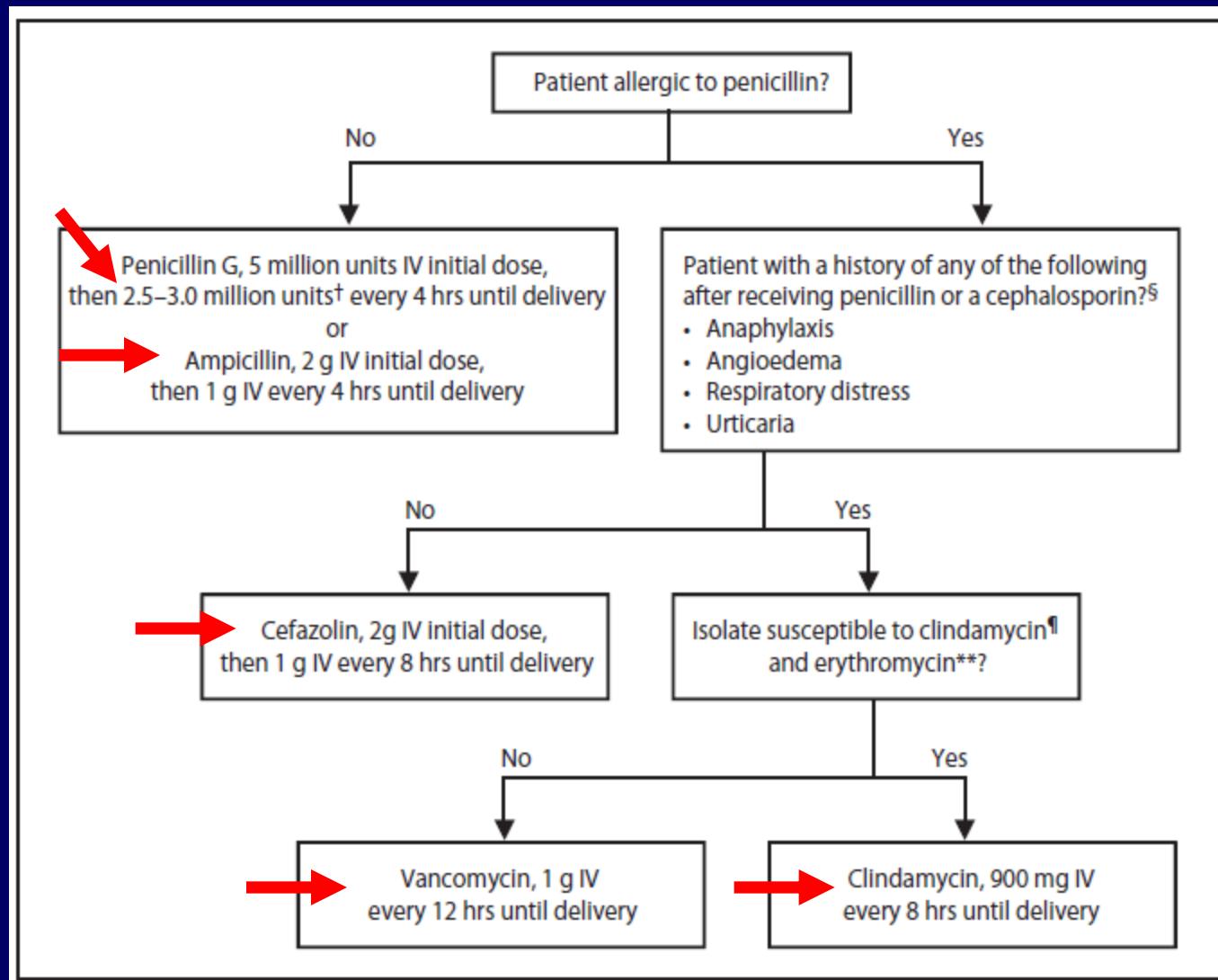


***S. agalactiae* versus erythromycin**  
***S. agalactiae* versus clindamycin**

**Percentage Susceptible**



# INTRAPARTUM PROPHYLAXIS



MMWR. 59 (RR-10): 1-32; 2010

# THE TIMES...

ANTIMICROBIAL AGENTS AND CHEMOTHERAPY, Aug. 2008, p. 2890–2897  
0066-4804/08/\$08.00+0 doi:10.1128/AAC.00185-08  
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Vol. 52, No. 8

## First Molecular Characterization of Group B Streptococci with Reduced Penicillin Susceptibility<sup>▽</sup>

Kouji Kimura,<sup>1</sup> Satowa Suzuki,<sup>1</sup> Jun-ichi Wachino,<sup>1</sup> Hiroshi Kurokawa,<sup>1</sup> Kunikazu Yamane,<sup>1</sup> Naohiro Shibata,<sup>1</sup> Noriyuki Nagano,<sup>1,2</sup> Haru Kato,<sup>1</sup> Keigo Shibayama,<sup>1</sup> and Yoshichika Arakawa<sup>1\*</sup>

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Received 9 February 2008/Returned for modification 8 March 2008/Accepted 5 May 2008



- Fourteen non-invasive *S. agalactiae* isolates between 1995-2005 had alterations in PBP2X
- Clinical significance unclear

Antimicrobial. Agents Chemother. 52: 2890-2897; 2008

# ...THEY ARE A CHANGIN'

J Antimicrob Chemother 2010  
doi:10.1093/jac/dkp458  
Advance publication 18 December 2009

## Prosthetic hip joint infection with a *Streptococcus agalactiae* isolate not susceptible to penicillin G and ceftriaxone

Christiane Gaudreau<sup>1,2\*</sup>, René Lecours<sup>3</sup>,  
Johanne Ismail<sup>4</sup>, Simon Gagnon<sup>5</sup>, Louise Jetté<sup>6</sup> and  
Michel Roger<sup>2,5</sup>

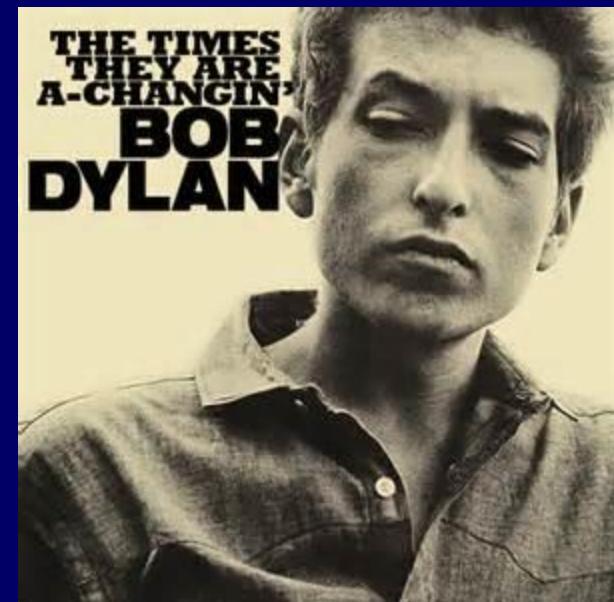
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**Table 1.** MICs (mg/L) of antimicrobial agents for GBS isolated in 2004 and 2007

	GBS 2004	GBS 2007	CLSI S	CLSI R
Penicillin G	0.06	0.25	≤0.12	NA
Ceftriaxone	0.12	1	≤0.5	NA
Oxacillin <sup>a</sup>	1	4	NA	NA
Ampicillin	0.12	0.5	≤0.25	NA
Meropenem	0.03	0.25	≤0.5	NA
Erythromycin	0.06	0.06	≤0.25	≥1
Clindamycin	0.06	0.12	≤0.25	≥1
Tetracycline	32	32	≤2	≥8
Vancomycin	0.25	0.5	≤1	NA
Levofloxacin	0.5	0.5	≤2	≥8
Chloramphenicol	4	4	≤4	≥16

GBS, group B *Streptococcus*; CLSI S and CLSI R, MIC breakpoints for susceptibility (S) and resistance (R);<sup>a</sup> NA, not available.

<sup>a</sup>No breakpoints for GBS, but breakpoints for *Staphylococcus* spp. are: ≥4 mg/L, resistant; and ≤2 mg/L, susceptible.



J. Antimicrob. Chemother. 65: 594-595; 2010

ANTIMICROBIAL AGENTS AND CHEMOTHERAPY, June 2011, p. 2983–2985  
0066-4804/11/\$12.00 doi:10.1128/AAC.01243-10  
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Vol. 55, No. 6

## Novel Mutations in a Patient Isolate of *Streptococcus agalactiae* with Reduced Penicillin Susceptibility Emerging after Long-Term Oral Suppressive Therapy<sup>▽</sup>

Jean Longtin,<sup>1,2\*</sup> Christie Vermeiren,<sup>3,4</sup> Dea Shahinas,<sup>1,4</sup> Gurdip Singh Tamber,<sup>1,4</sup> Allison McGeer,<sup>4,5</sup> Donald E. Low,<sup>1,4,5</sup> Kevin Katz,<sup>3,4,6</sup> and Dylan R. Pillai<sup>1,4</sup>

Ontario Agency for Health Protection and Promotion, Toronto, Ontario, Canada<sup>1</sup>; Centre Hospitalier Universitaire de Québec, Québec, Canada<sup>2</sup>; Shared Hospital Laboratory, Toronto, Ontario, Canada<sup>3</sup>; University of Toronto, Toronto, Ontario, Canada<sup>4</sup>; Mount Sinai Hospital, Toronto, Ontario, Canada<sup>5</sup>; and North York General Hospital, Toronto, Ontario, Canada<sup>6</sup>

# *S. agalactiae* BACTERIURIA

- Marker for heavy genital tract colonization; risk factor for early-onset GBS disease  
Scand. J. Infect. Dis. 17: 195-199; 1985
- 1996 guidelines      no threshold specification  
2002 guidelines      report any concentration  
2010 guidelines       $10^4$  colony forming units/mL
- Few data available on risk for early-onset GBS in context of low-count bacteriuria  
MMWR. 45 (RR-7): 1-24; 1996  
MMWR. 51 (RR-11): 1-24; 2002  
MMWR. 59 (RR-10): 1-32; 2010

# THE END

- Identification of candidates for intrapartum chemoprophylaxis is essential for prevention of early-onset group B streptococcal disease
- Much of this has fallen into the hands of laboratory
- Situation has improved since the 1970s; more work to be done
- Molecular diagnostics and antimicrobial susceptibility testing, when applied appropriately, play major role

