

MRE (multiresistente Erreger)

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MRE

- MRSA, VRE
- ESBL+ (sonstige multiresistente gramnegative Bakterien)
- MDR-/XDR-Tuberkulose

MRE

- MRSA, VRE
- ESBL+ (sonstige multiresistente gramnegative Bakterien)
- MDR-/XDR-Tuberkulose

MRSA

MRSA

- M=Methicillin (=Oxacillin):
 - bedeutet Resistenz gegenüber β -Lactame
- viele MRSA-Stämme sind (hier) gut empfindlich auf ältere Antibiotika, z.B.
 - Doxycyclin
 - Cotrimoxazol
- es gibt einige neue MRSA-wirksame Antibiotika (incl. β -Lactame !!!)
- (Screening +) Dekolonisierung möglich

MRSA-Dekolonisierung

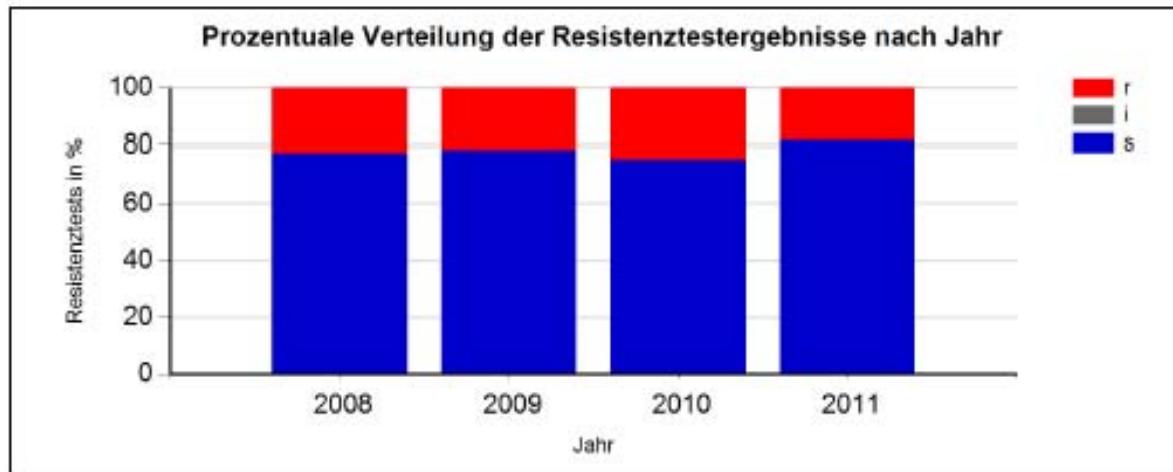
- Evaluation der holländischen Strategie seit 2006
- Mupirocin + Chlorhexidin (+ systemische Antibiotika bei “komplizierten” Fällen)
- N=613
- Erfolg 60% nach einmaligem Zyklus, 80% insgesamt
- Behandlung nach Protokoll (natürlich) erfolgreicher

Parameter:

Erreger:	<i>Staphylococcus aureus</i>	Materialgruppe:	Blutkultur
Antibiotikum:	Oxacillin	Fachrichtung:	Alle
Versorgungsbereich:	stationäre Versorgung	Stationstyp:	Alle
Intervall:	Jahr	Versorgungsstufe:	Alle

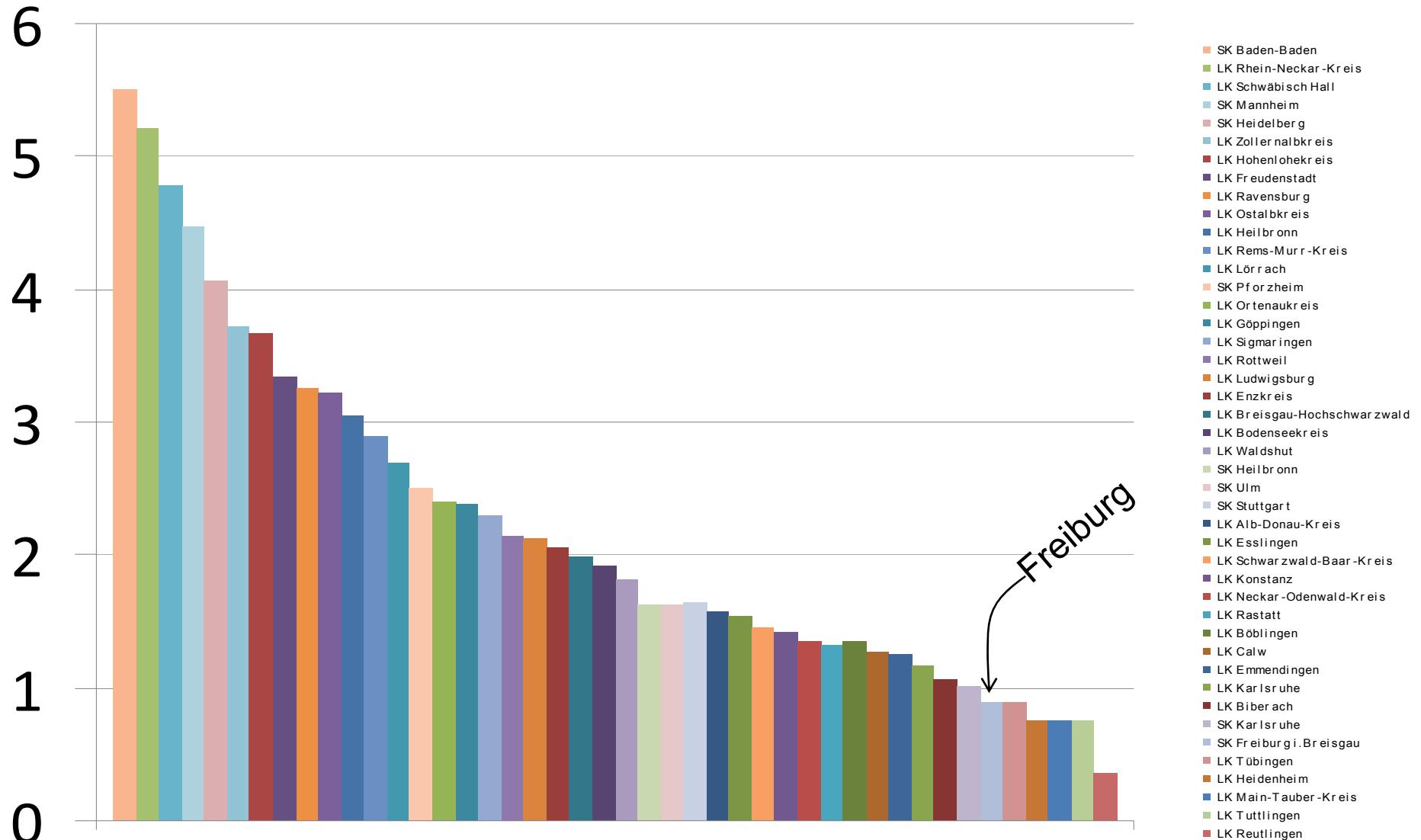
Datenstand: 26.06.2012

Intervall	R		I		S		Total
	n	%	n	%	n	%	
Jahr 2011	296	18,3	0	0,0	1324	81,7	1620
Jahr 2010	363	24,7	0	0,0	1104	75,3	1467
Jahr 2009	271	21,4	0	0,0	998	78,6	1269
Jahr 2008	177	22,3	0	0,0	615	77,7	792



MRSA-“Sepsis”-Inzidenz 2011

(pro 100.000 Einwohner und Jahr; Quelle: SurvStat@RKI 7.3.2012)



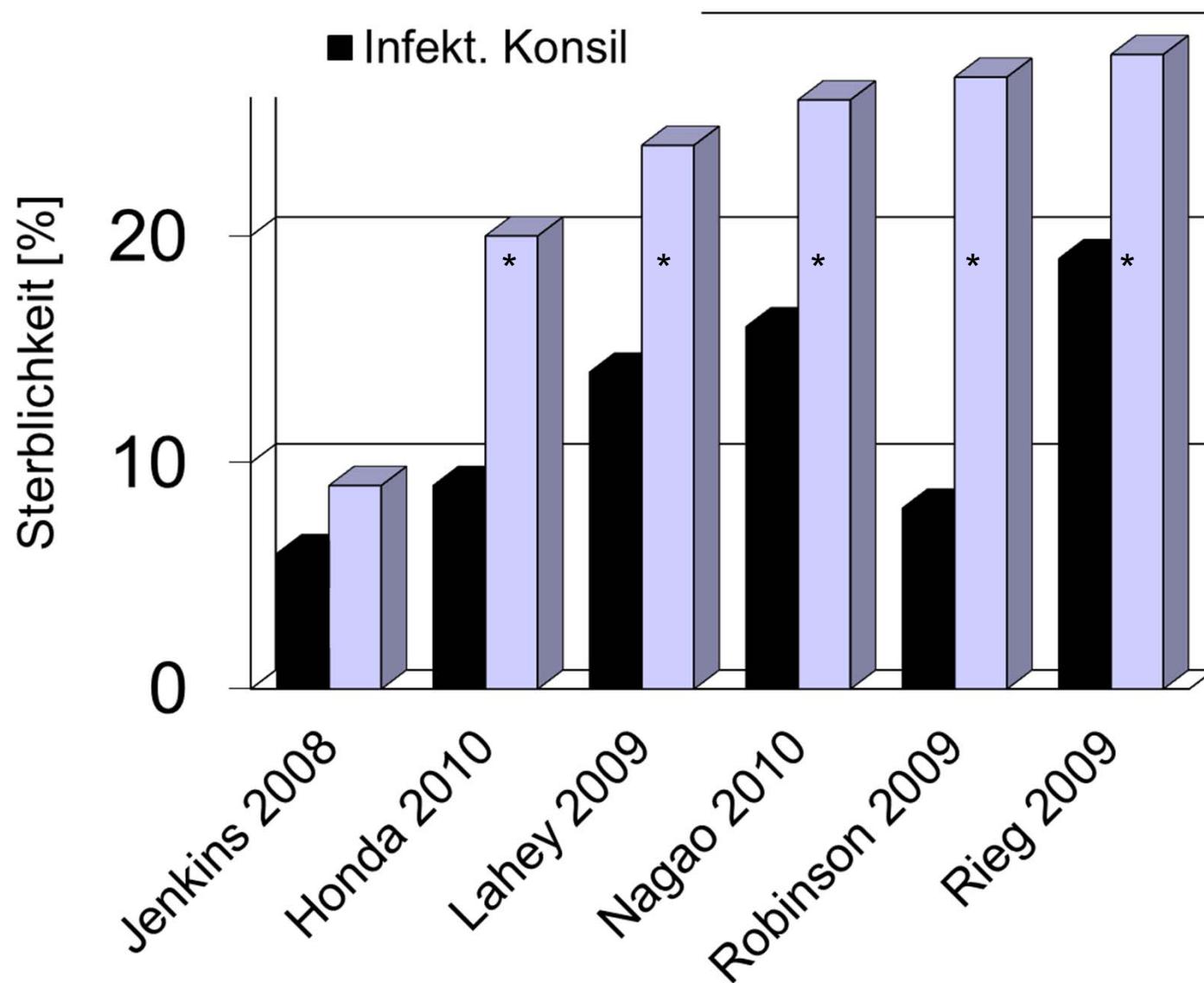
MRSA-Infektion

MRSA-Infektion

- Haut-Weichteil- und Wundinfektionen
- Knochen- und Gelenkinfektion
- Pneumonie
- Bakterämie/Sepsis
 - Unkompliziert
 - Kompliziert
 - Verschiedne Foci/Lokalisationen



S. aureus-Bakterämie (SAB)



Therapie der MRSA-Infektion

- „traditionell“: Vancomycin

MRSA

- viele MRSA-Stämme sind (hier) gut empfindlich auf ältere Antibiotika, z.B.
 - Doxycyclin
 - Cotrimoxazol
- es gibt einige neue MRSA-wirksame Antibiotika (incl. β -Lactame !!!)

MRSA: Therapieoptionen (in Deutschland)

Tab. 4.1.2.2: Resistenz gegen Antibiotika (zusätzlich zur Resistenz gegen β -Lactamantibiotika) bei ha-MRSA, 2006–2010

Antibiotikum	2006 (%)	2007 (%)	2008 (%)	2009 (%)	2010 (%)
Oxacillin	100	100	100	100	100
Ciprofloxacin	93,8	95,8	91	90	86
Moxifloxacin	96,3	94,4	89,6	87	86
Erythromycin	72,5	75	80,7	67	65
Clindamycin	65,4	72	73,4	60	59
Gentamicin	13,3	9,8	10,5	9,5	5,3
Oxytetracyclin	7,4	6,8	7,3	8	6,0
Rifampicin	2,5	1,07	0,4	1,6	0,8
Cotrimoxazol	3,1	2	10,8	5,3	0,8
Fusidinsäure	6,4	3,8	2,0	5,2	4,0
Fosfomycin	3,3	0,56	1,1	0,15	0,6
Linezolid	0,04	0,11	0,1	0,1	0,08
Tigecyclin	0	0	0	0	0,12
Daptomycin	0	0	0,65	1,3	1,6
Mupirocin	2,6	3,3	5,3	4,0	4,6
Vancomycin	0	0	0	0	0,08
Telcoplanin	0	0	0	0	0,2

MRSA: Therapieoptionen ?

■ MRSAB/schwere Infektion

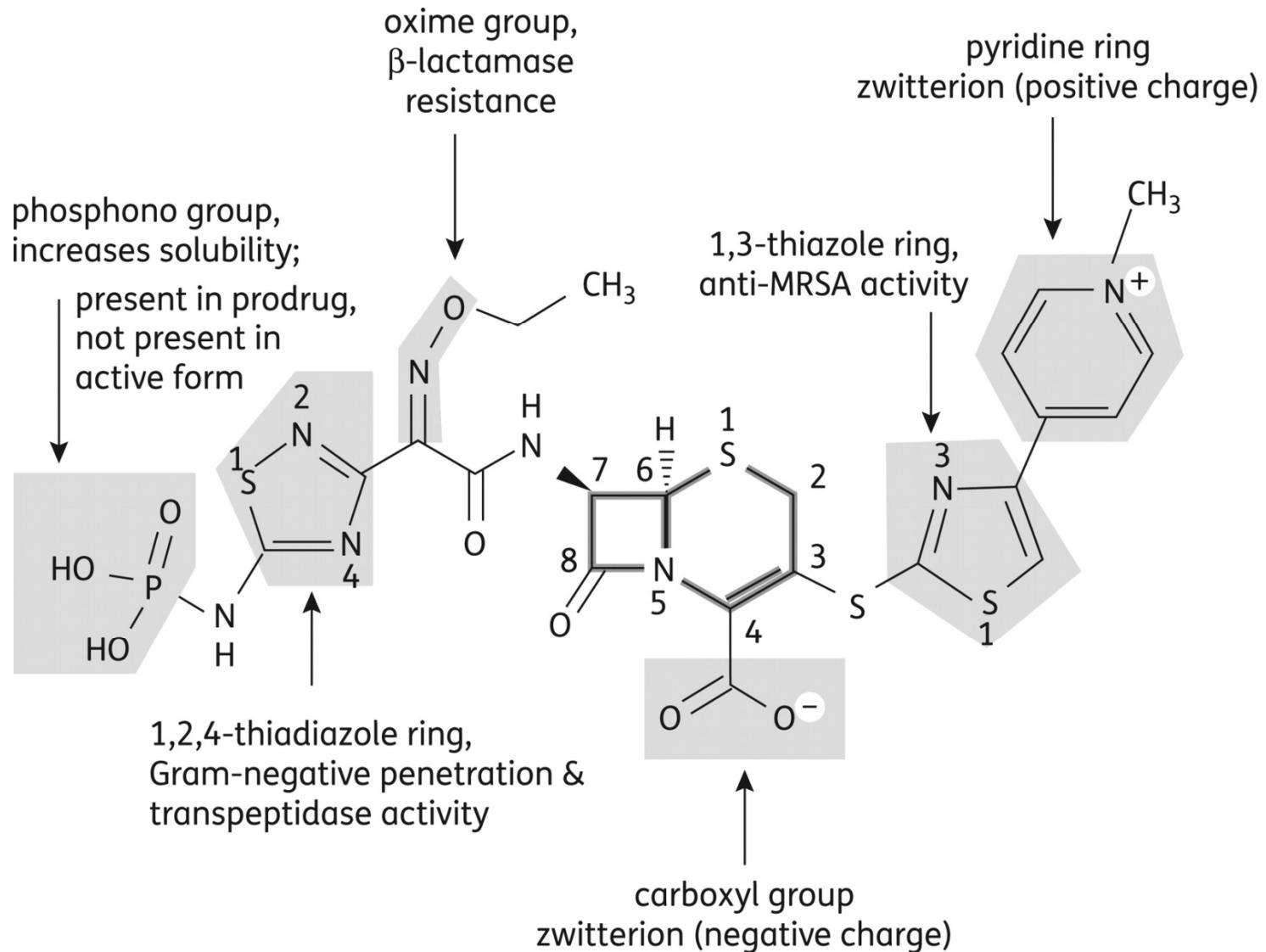
- Vancomycin-Dosisescalation ☹
- Kombination, z.B. Rifampicin ☺?
- alternative Substanzen ☺
 - Daptomycin (ok, nicht sicher besser) (nicht: Pneumonie)
 - Linezolid (bei Pneumonie besser und ok, aber bakteriostatisch)
 - sonst: wenig Daten

■ SSTI und andere lokalisierte Infektionen (cave: Fremdkörper)

- ?
- Doxycyclin, Citrimoxazol, Linezolid, Clindamycin
- Fusidinsäure, Pristinamycin, ...

... neue MRSA- β -Lactame ??

Ceftarolin-Fosamil (Prodrug)



Ceftarolin

- MHK₉₀ ($\mu\text{g}/\text{mL}$) Staphylokokken
 - MSSA 0.25
 - MRSA 1.0

- MHK₉₀ ($\mu\text{g}/\text{mL}$) Pneumokokken
 - PenS 0.015
 - PenR 0.125

Ceftarolin (u.a.)

- (gut) studiert bei (und zugelassen für)
 - ambulant erworbene Pneumonie
 - Haut-Weichteilinfektionen
- Kaum Erfahrung bei schweren MRSA-Infektionen, hier möglicherweise unterdosiert (?)





Analyse von Fleischproben auf MRSA und ESBL-produzierende Keime – Fragen und Antworten

Autorin: Dr. Kathrin Birkel

Stand: 24. Januar 2012

Quellen für alle Angaben liegen beim BUND vor.

BUND-Stichprobenuntersuchung von Hähnchenfleisch auf MRSA und ESBL-produzierende Keime - Befunde Dezember 2011			
Ort/ Einkaufsstätte	Marke	Produktionsfirma	Befund
Berlin			
Edeka	Heidegold	Wiesenhof	ESBL: positiv - E.coli; MRSA: negativ
Edeka	Heidegold	Wiesenhof	ESBL: negativ; MRSA: negativ
Penny	Juwel	Stolle	ESBL: positiv - E.coli; MRSA: negativ
Penny	Juwel	Stolle	ESBL: positiv - E.coli; MRSA: negativ
Hamburg			
Netto	Gut Ponholz	Stolle	ESBL: positiv - E.coli; MRSA: negativ
Netto	Gut Ponholz	Stolle	ESBL: positiv MRSA: negativ
Rewe	Wilhelm Brandenburg	Sprehe	
Rewe	Wilhelm Brandenburg	Sprehe	
Köln			
Netto			

ESBL positiv 10/20

Aldi	geka	Wiesenhof	ESBL: negativ; MRSA: negativ
Aldi	geka	Wiesenhof	ESBL: negativ; MRSA: negativ
Edeka	Astenhof	Sprehe	ESBL: positiv - E.coli; MRSA negativ
Edeka	Astenhof	Sprehe	ESBL: negativ; MRSA: negativ
Stuttgarter Region			
Aldi	Landgeflügel	Rothköter	ESBL: negativ; MRSA: negativ
Metro	Astenhof	Sprehe	ESBL: negativ; MRSA: negativ
Lidl	Landjunker	Wiesenhof	ESBL: positiv - E.coli; MRSA: negativ
Kaufland	Wiesenhof	Wiesenhof	ESBL: negativ; MRSA: negativ

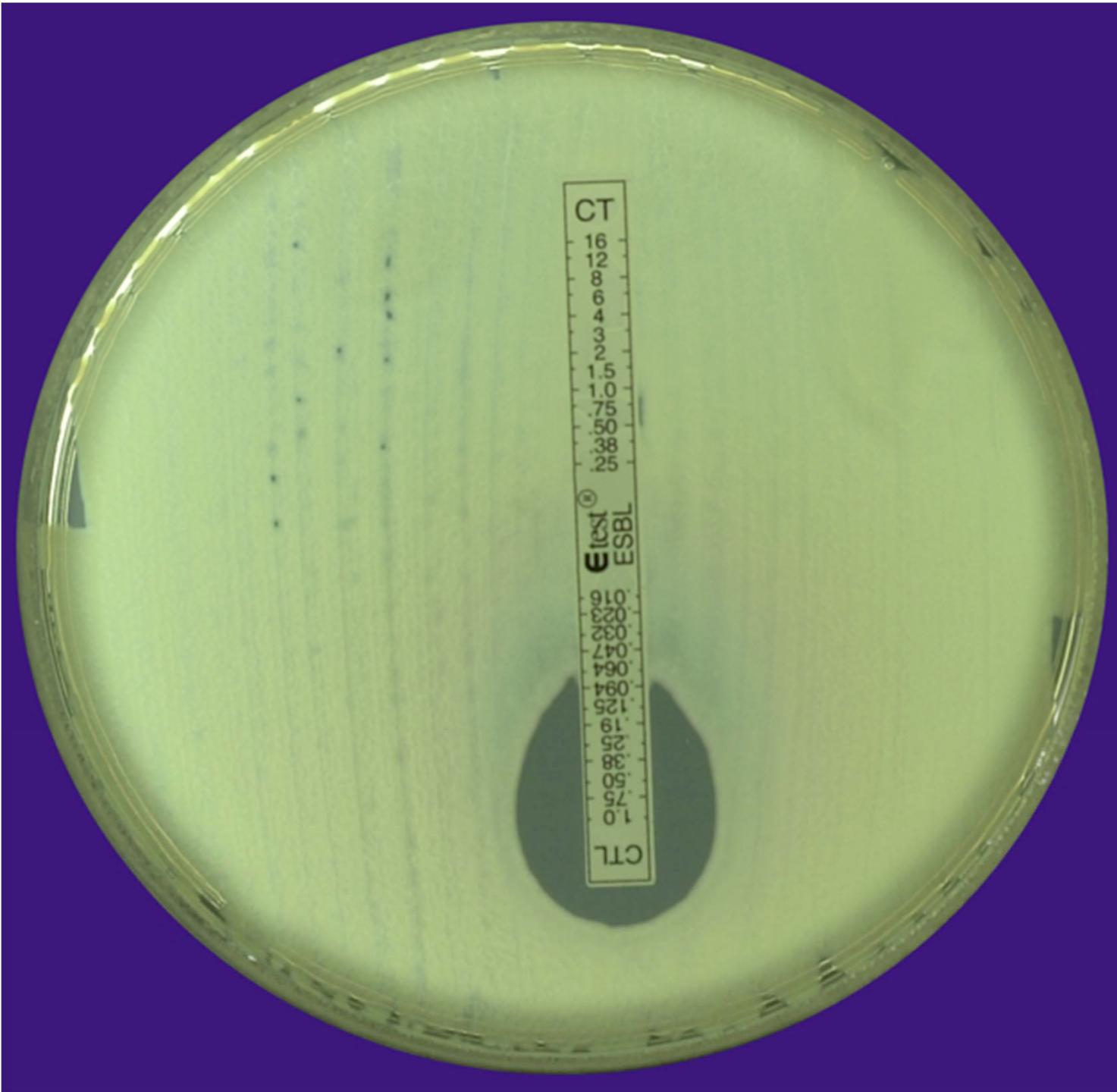
... a total of 399 chicken meat samples from nine supermarket chains, four organic food stores and one butcher's shop in two geographically distinct regions (Berlin and Greifswald) were screened for ESBL production ...

.... ESBL isolates were obtained from 175 samples (43.9%) from all tested sources.

No differences could be observed in the prevalence of ESBL producers between organic and conventional samples.

73% of the ESBL producers showed co-resistance to tetracycline, 36% to co-trimoxazole and 8% to ciprofloxacin





From MICs to S / I / R : where are the limits ?

Which breakpoint do you use ?

before ...



now ...
(2009)

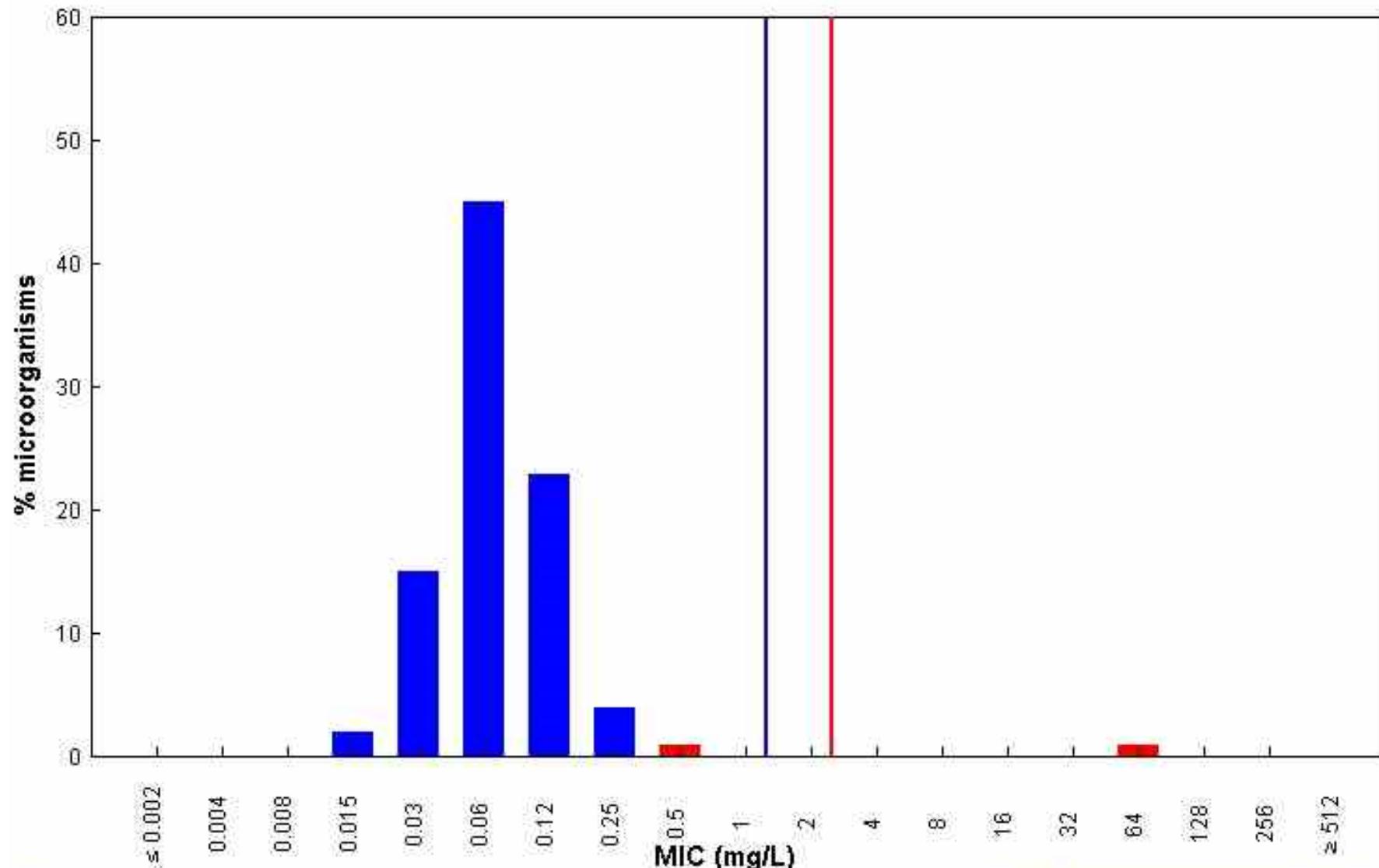


cefotaxime vs. <i>E.coli</i>	\leq S / R
BSAC United Kingdom	2 / \geq 4
CA-SFM France	4 / > 32
CRG The Netherlands	4 / > 16
DIN Germany	2 / \geq 16
NWGA Norway	1 / \geq 32
SRGA Sweden	0.5 / \geq 2

cefotaxime vs. <i>E.coli</i>	\leq S / R
EUCAST EU	1 / 2
CLSI USA	8 / \geq 64

Cefotaxime / Escherichia coli
EUCAST MIC Distribution - Reference Database 2013-03-06

MIC distributions include collated data from multiple sources, geographical areas and time periods and can never be used to infer rates of resistance



MIC

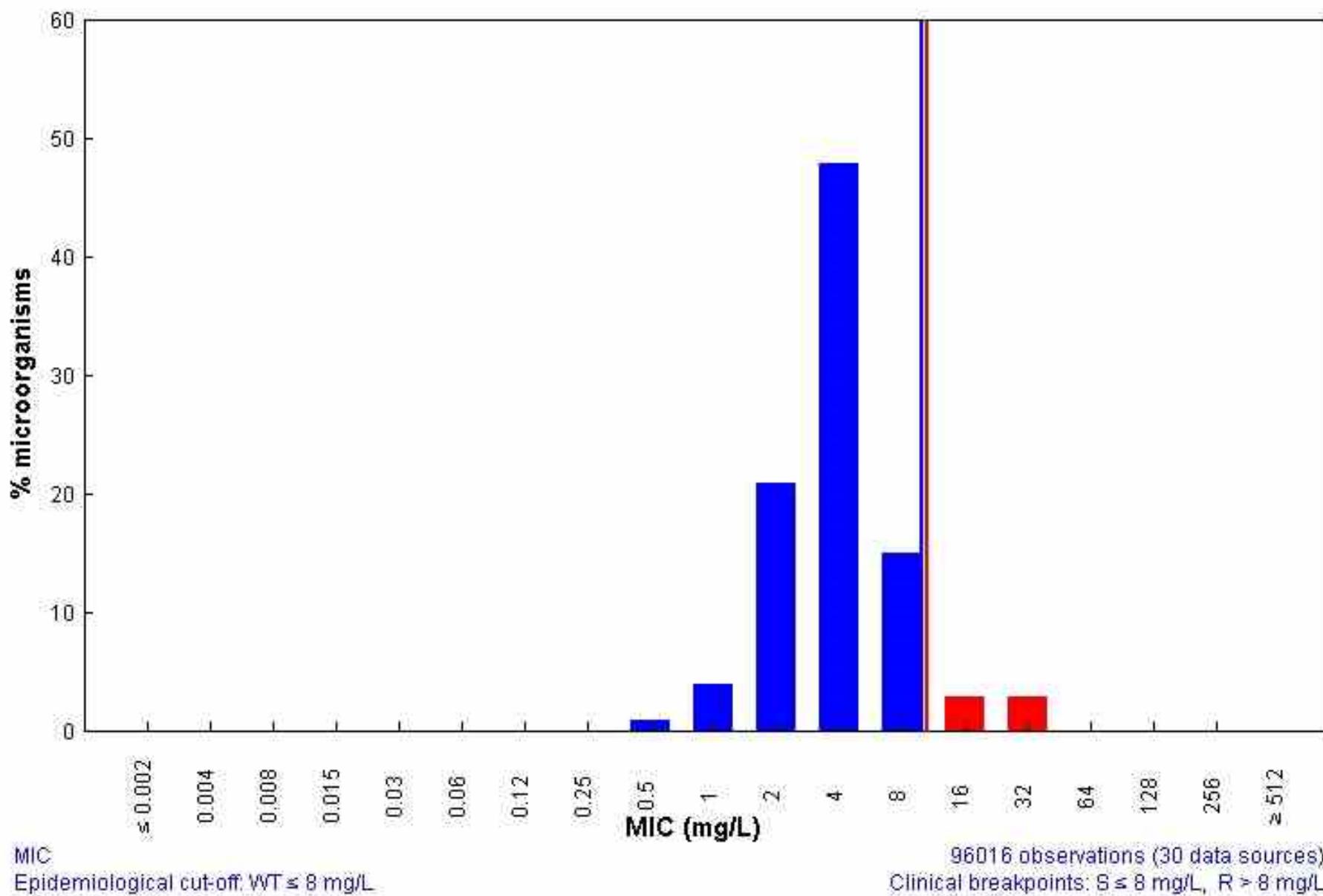
Epidemiological cut-off: WT ≤ 0.25 mg/L

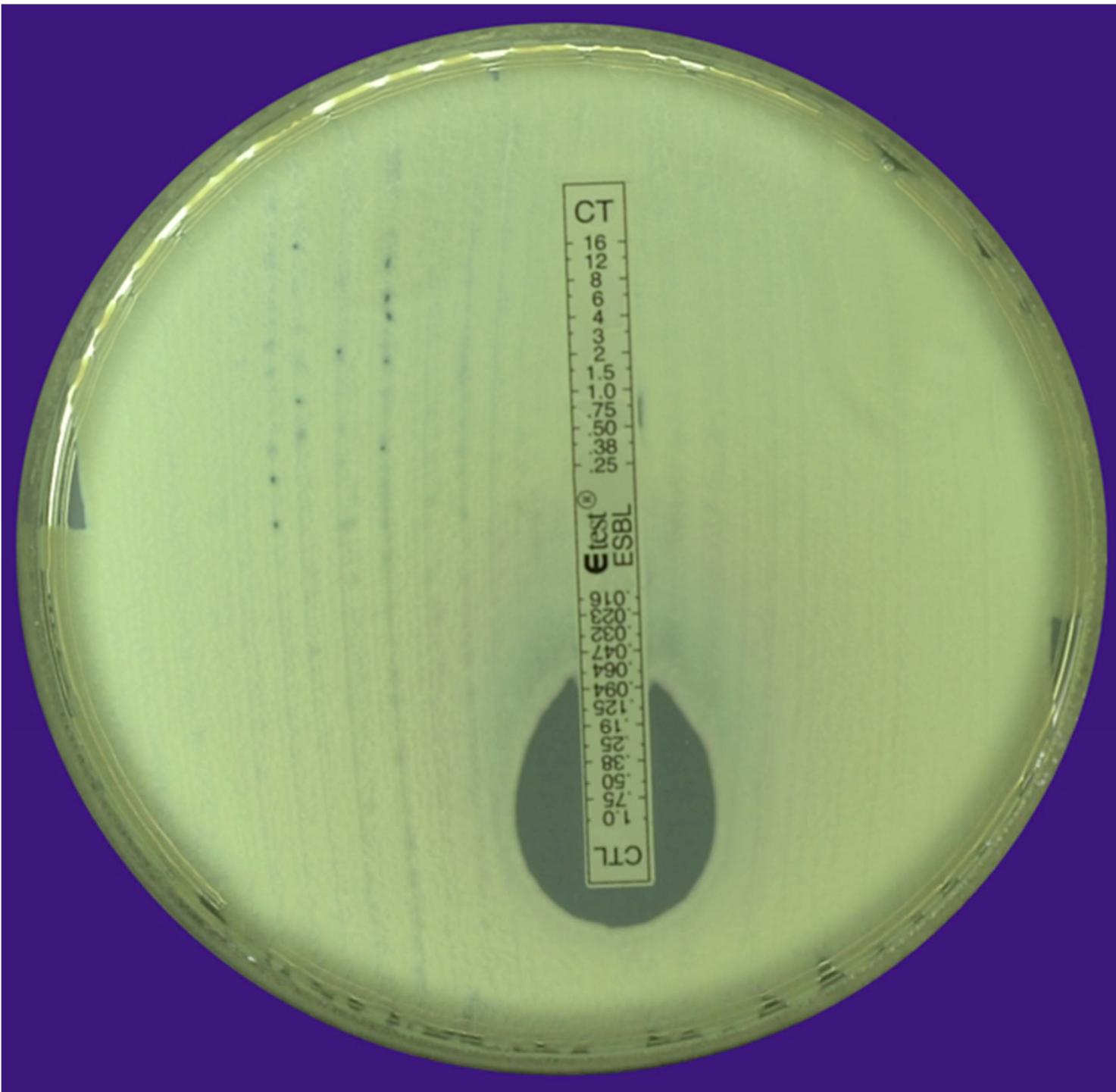
10855 observations (58 data sources)

Clinical breakpoints: S ≤ 1 mg/L, R > 2 mg/L

Cefuroxime / Escherichia coli
EUCAST MIC Distribution - Reference Database 2013-03-06

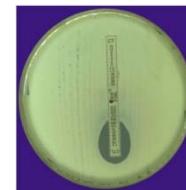
MIC distributions include collated data from multiple sources, geographical areas and time periods and can never be used to infer rates of resistance





ESBL

- gelten als resistent gegenüber den meisten Penicillinen und Cephalosporinen (enzymatische Spaltung = *extended spectrum β-lactamase*, verschiedene Typen)
 - Cefotaxim/Ceftriaxon manchmal unterschiedlich zu Ceftazidim, oft unterschiedlich zu Cefepim
 - Starker Inokulum-Effekt
 - Ceph III° + Clav meistens aktiv



Detection of Favorable Oral Cephalosporin-Clavulanate Interactions by *In Vitro* Disk Approximation Susceptibility Testing of Extended-Spectrum-Beta-Lactamase-Producing Members of the *Enterobacteriaceae*

Jennifer D. Campbell,^{a,b,c*} James S. Lewis II,^{b,c,e} M. Leticia McElmeel,^d Letitia C. Fulcher,^d and James H. Jorgensen^d

TABLE 2 Results of tests of cefpodoxime with and without the concomitant presence of amoxicillin-clavulanate (Amox-Clav) by broth dilution and disk diffusion testing

ESBL group (no. of isolates)	MIC tested alone						Cefpodoxime MIC tested with Amox-Clav ^a						% susceptible ^b	% detected by disk approximation ^c		
	Cefpodoxime			Amox-Clav ^d			Cefpodoxime MIC tested with Amox-Clav ^a									
	MIC ₅₀ ^e	MIC ₉₀ ^f	Range	MIC ₅₀	MIC ₉₀	Range	MIC ₅₀	MIC ₉₀	Range							
CTX-M (28) ^g	>32	>32	16->32	16	32	4->32	0.25	8	≤0.06->32	89.3			100			
SHV (17) ^h	>32	>32	2->32	32	>32	4->32	0.5	>32	≤0.06->32	58.8			100			
CTX-M + SHV (3) ⁱ	>32	>32	>32	32	>32	16->32	1	>32	≤0.06->32	66.7			100			

ESBL

- Amoxi+Clav (Amp+Sul) und Pip+Tazo können aktiv sein
- häufig auch resistent gegenüber
 - Fluorchinolone
 - Aminoglykoside
 - Cotrimoxazol

ESBL

- meist empfindlich gegenüber
 - Fosfomycin
 - Nitrofurantoin
 - Pivmecillinam
 - Nitroxolin
- Carbapeneme bei schweren Infektionen als Initialtherapie derzeit Mittel der Wahl

ESBL Trends weltweit

	2003	2004	2005	2006	2007
Asia/Pacific	15%	14%	23%	28%	40%
South America	10%	12%	18%	23%	30%
Middle East/ Africa	8%	11%	16%	17%	17%
Europe	6%	3%	8%	9%	10%
North America	4%	1%	4%	8%	8%

Source: SMART-Programme (Study for Monitoring Antimicrobial Resistance Trends) with collection of *E. coli*, *Klebsiella pneumoniae* und *K. oxytoca* from 74-94 laboratories since 2003

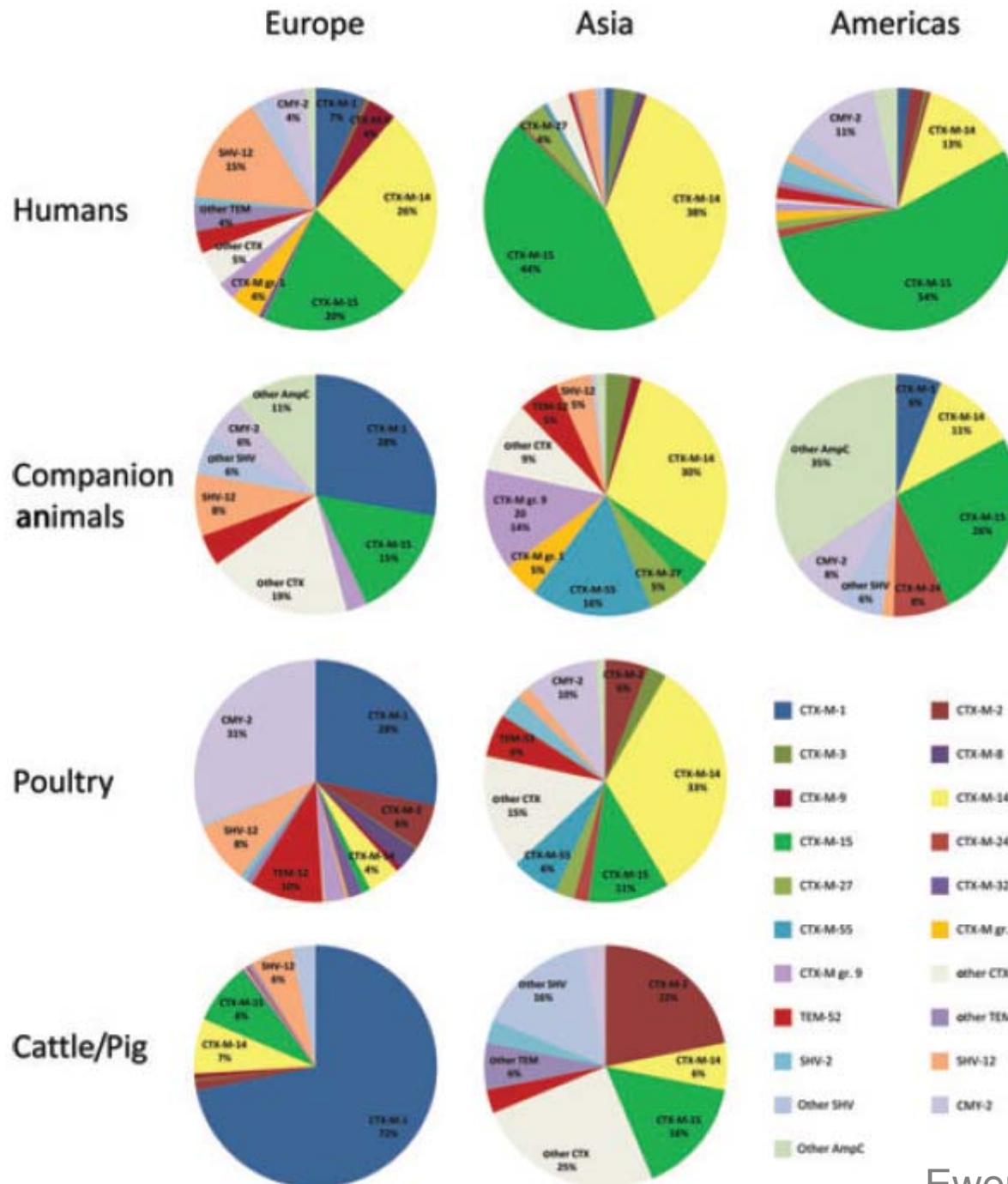
ESBL Trends Asien/Pazifikraum

- 2008, SMART
- Gram-negative bacilli associated with intra-abdominal infections
- CLSI
- ESBL+ *E. coli* + *Klebsiella pneumoniae*

China	59%	34%
India	61%	47%
Thailand	53%	23%
- FQREC

Thailand	>50%
China	≥70%
India	>80%

ESBL+/ AmpC+ *E. coli*



Ewers et al. CMI 2012

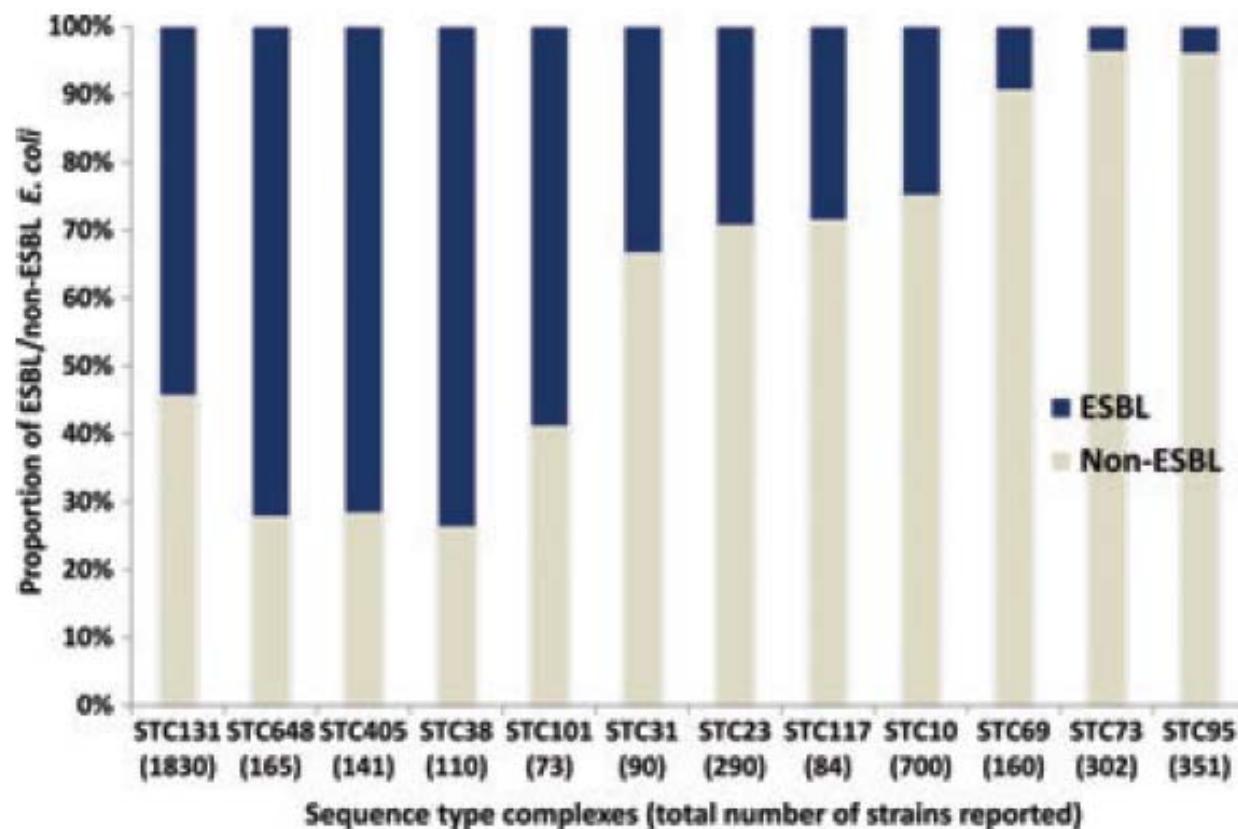


FIG. 4. Proportion (%) of extended-spectrum β -lactamase (ESBL)/AmpC/NDM-producing *Escherichia coli* among the total number of strains recorded among various sequence type complexes (STCs).

ESBL-“Erwerb“

- 242 Patienten mit Reisediarrhoe
- ESBL+ *E. coli*, *Klebsiella pneumoniae*,
Proteus mirabilis bei 24% (58/242)
- Europa, 3% (2/63) vs RoW 36% (50/138)
 - Indien: 11/14 (79%)
 - Ägypten 19/38 (50%)
 - Thailand 8/38 (22%)

ESBL Trends Afrika, z.B.

- Kamerun 2009
- Stuhlproben von 208 Patienten + 150 Gesunden
- ESBL+ 23% vs 7%
- Risikofaktor: Fluorchinolon-Vorbehandlung

ESBL-Übertragung

- Adoptivkinder aus Mali (→Frankreich)
- Stuhlproben bei Ankunft sowie später in monatlichen Abständen
- Stuhlproben der Adoptiveltern und Familienmitglieder
- Ergebnisse
 - ESBL bei 24/25 Adoptivkindern nachgewiesen
 - mehrheitlich *E. coli*, aber auch *Salmonella*
 - Kolonisationsdauer im Mittel 9 Monate
 - Übertragung intrafamiliär in 5/22 Familien (23%)

Ceph III°-resistente *E.coli* hier

stationär: 10%
ambulant: 5%

E.coli - alle Mat. – IM/Allg vs. Pädiater vs. Urologen vs stationär

<i>E.coli</i>	R (%) Innere/Allg	R (%) Pädiater	R (%) Urologen	R (%) stationär
Amoxi/Clav	22	15	24	31
Cefotaxim	5	3	6	10
FQ	18	4	24	22
Cotrim	29	22	32	31

ESBL: Optionen ?

- Carbapenem
- ?? einige sind in vitro empfindlich auf Amoxi/Clav bzw. Pip/Tazo

β -Lactam/ β -Lactam Inhibitor Combinations for the Treatment of Bacteremia Due to Extended-Spectrum β -Lactamase-Producing *Escherichia coli*: A Post Hoc Analysis of Prospective Cohorts

Jesús Rodríguez-Baño,^{1,2} María Dolores Navarro,¹ Pilar Retamar,¹ Encarnación Picón,¹ Álvaro Pascual,^{1,3} and the Extended-Spectrum Beta-Lactamases–Red Española de Investigación en Patología Infecciosa/Grupo de Estudio de Infección Hospitalaria Group^a

Characteristic	Empirical Therapy Cohort		
	BLBLI (n = 72)	Carbapenem (n = 31)	P
Age, median y (IQR)	69 (59–80)	60 (52–78)	.1 ^b
Male sex	29 (40.3)	11 (35.5)	.6
Nosocomial acquisition	26 (36.1)	24 (77.4)	<.001
Charlson index, median, (IQR)	2 (1–5)	2 (1–5)	.6 ^b
Cancer	21 (31.9)	11 (35.5)	.7
Immunosuppression	5 (6.9)	5 (16.1)	.1 ^c
Neutropenia	2 (2.8)	3 (9.7)	.1 ^c
Urinary or biliary tract as source	52 (72.2)	18 (58.1)	.1
ICU admission	7 (9.9)	2 (6.7)	.7 ^c
Severe sepsis or shock at presentation	14 (19.4)	9 (29.0)	.2
Pitt score, median (IQR)	1 (0–2)	1 (0–2)	.7 ^b
CTX-M enzyme	57 (80.3)	25 (86.2)	.4
Definitive therapy			
Carbapenem	32 (44.4)	30 (93.7)	<.001
BLBLI	34 ^d (47.2)	0	<.001
Empirical therapy			
Carbapenem
BLBLI
Cephalosporins
Fluoroquinolones
Appropriate empirical therapy
Mortality, no. of deaths			
Day 7	2 (2.8)	3 (9.7)	.1 ^c
Day 14	7 (9.7)	5 (16.1)	.3
Day 30	7 (9.7)	6 (19.4)	.1
Hospital stay after BSI , median (IQR), d	12 (8–28)	13 (9–25)	.7 ^b

Characteristic	Definitive Therapy Cohort		
	BLBLI (n = 54)	Carbapenem (n = 120)	P
Age, median y (IQR)	67 (56–83)	70 (55–78)	.3 ^b
Male sex	34 (63)	70 (58.3)	.5
Nosocomial acquisition	18 (33.3)	67 (55.8)	.006
Charlson index, median, (IQR)	2.5 (1–5)	3 (1–5)	.5 ^b
Cancer	15 (27.8)	43 (35.8)	.2
Immunosuppression	3 (5.6)	15 (12.5)	.1
Neutropenia	0	7 (5.8)	.1 ^c
Urinary or biliary tract as source	42 (77.8)	79 (65.8)	.1
ICU admission	4 (7.4)	18 (15.4)	.1
Severe sepsis or shock at presentation	8 (14.8)	32 (26.7)	.08
Pitt score, median (IQR)	1 (0–2)	1 (1–2)	.04 ^b
CTX-M enzyme	43 (82.7)	95 (81.2)	.8
Definitive therapy			
Carbapenem
BLBLI
Empirical therapy			
Carbapenem	0	30 (25)	<.001
BLBLI	45 ^d (83.3)	38 (31.7)	<.001
Cephalosporins	7 (13)	39 (32.5)	.006
Fluoroquinolones	2 (3.7)	13 (10.8)	.1 ^c
Appropriate empirical therapy	34 (63)	64 (53.3)	.2
Mortality, no. of deaths			
Day 7	1 (1.9)	5 (4.2)	.6 ^c
Day 14	3 (5.6)	14 (11.7)	.2
Day 30	5 (9.3)	20 (16.7)	.1
Hospital stay after BSI , median (IQR), d	13 (8–22)	13 (10–25)	.04 ^b

Table 4. Cox Regression Analysis of Associations Between Different Variables and Mortality in the Definitive Therapy Cohort

Characteristic	Definitive Therapy Cohort		
	BLBLI (n = 54)	Carbapenem (n = 120)	P
Age, median y (IQR)	67 (56–83)	70 (55–78)	.3 ^b
Male sex	34 (63)	70 (58.3)	.5
Nosocomial acquisition	18 (33.3)	67 (55.8)	.006
Charlson index, median, (IQR)	2.5 (1–5)		
Cancer			
Variable	Crude Analysis		Adjusted Analysis
	HR (95% CI)	P	HR (95% CI) P
Male sex	1.2 (46–2.29)	.9	...
Age ^a	1.00 (97–1.02)	.9	...
Nosocomial BSI	0.99 (45–2.22)	.9	...
Charlson index ^a	1.02 (88–1.28)	.7	...
Neutropenia	1.78 (88–13.32)	.5	...
High-risk source ^b	2.07 (94–4.54)	.06	...
Pitt score ^a	1.49 (1.26–1.78)	<.001	1.38 (1.12–1.70) .002
Severe sepsis or shock ^c	2.07 (94–4.54)	.001	2.10 (87–5.05) .09
Empirical therapy with BLBLI	3.64 (1.66–7.99)	.3	...
Inappropriate empirical therapy	0.56 (1.18–1.73)	.1	...
Definitive therapy with BLBLI ^d	1.76 (78–3.93)	.4	0.76 (28–2.07) .5
Mortality, no. of deaths			
Day 7	1 (1.9)	5 (4.2)	.6 ^c
Day 14	3 (5.6)	14 (11.7)	.2
Day 30	5 (9.3)	20 (16.7)	.1
Hospital stay after BSI , median (IQR), d	13 (8–22)	13 (10–25)	.04 ^b

Abbreviations: BLBLI, β -lactam/ β -lactamase inhibitor association; BSI, bloodstream infection; CI, confidence interval; HR, hazard ratio.

^a Charlson index, ^b High-risk source, ^c Pitt score, ^d Empirical therapy

ESBL: Optionen ?

- Carbapenem
- ! Amoxi/Clav oder Pip/Tazo (nach Austestung)

ESBL: Optionen ?

- Carbapenem
- ! Amoxi/Clav oder Pip/Tazo (nach Austestung)
- ?? Cephalosporin + β LI

Orthodox and unorthodox clavulanate combinations against extended-spectrum β -lactamase producers

D. M. Livermore, R. Hope, S. Mushtaq and M. Warner

Antibiotic Resistance Monitoring and Reference Laboratory, Health Protection Agency, Centre for Infections, London, UK

possible to launch a drug combination in India—a fast-developing country with one-sixth of the world's population—on the basis of what, in the West, would count as phase II data. Ceftriaxone-sulbactam is marketed on this basis by at least one local company (Ceftrimax, VHB Group, Mumbai, <http://www.vhbgroup.com>). Whilst, in the period between the Venice meeting and publication of this supplement, Ranbaxy (<http://www.ranbaxy.com>) have launched a cefepime-tazobactum combination. It will be intriguing to see the clinical results against ESBL producers and, if these are positive, the reactions of both western microbiologists and companies!

ESBL: Optionen ?

- Carbapenem
- ! Amoxi/Clav oder Pip/Tazo (nach Austestung)
- ? Cephalosporin + β LI
-
- ? Tigecycline *iv*
- ? Colistin *iv*
- Azithromycin (bei *Salmonella*)

ESBL: Optionen orale Therapie ?

- Orale Therapie (UTI)

- | | | |
|------------------------------|-----|--------------------------|
| ■ ? Fosfomycin <i>po</i> | 95% | in vitro-Empfindlichkeit |
| ■ ? Nitrofurantoin <i>po</i> | 90% | |
| ■ ? Nitroxolin <i>po</i> | 90% | |
| ■ ? Pivmecillinam <i>po</i> | 85% | |
| ■ ? Cefpodoxim+(Amoxi-)Clav | | |



SELEXID®

pivmecillinam

Selexid-Filmtabletten. **Zulassungsinhaber:** LEO Pharma, Wien. **Hersteller:** LEO Pharmaceutical Products, Ballerup, Danemark. **Zusammensetzung:** 1 Filmtablette enthält 200 mg Pivmecillinam-hydrochlorid. **Anwendungsgebiete:** Selexid kann bei Infektionen mit Mecillinam-empfindlichen Erregern eingesetzt werden, wie bei: Infektionen der Harnwege: akute Cystitis, Pyelitis, Pyelonephritis; chronische oder rezidivierende Harnwegsinfekte; postoperative Bakteriurie nach urologischen und gynäkologischen Eingriffen; symptomatische oder asymptomatische Bakteriurie in der Schwangerschaft; Salmonellosen, Shigellosen; als Alternative zur antibiotischen Behandlung des Typhus abdominalis; zur Sanierung von Salmonellenausscheidern; Enteritiden durch andere Mecillinam-empfindliche Bakterien

2 x 2 Tabl. à 200 mg 3 Tage

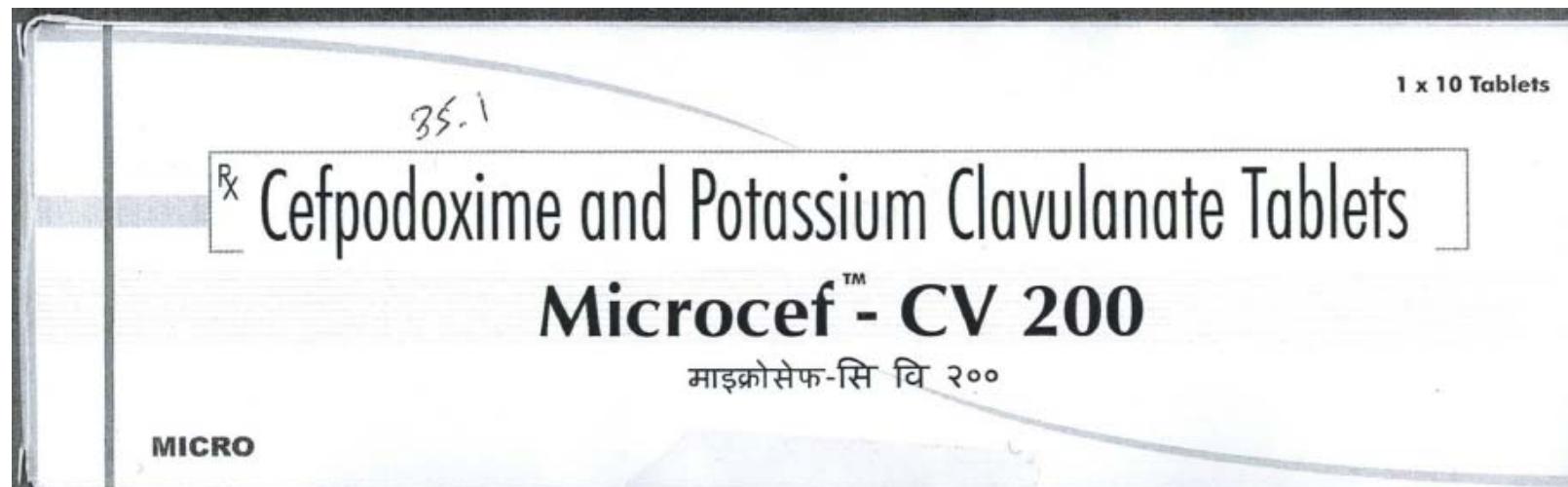
Spektrum und MHK-Werte (mg/l):

Keim	MHK ₅₀ (mg/l)	MHK ₉₀ (mg/l)
<i>Staphylococcus saprophyticus</i>	16 ¹⁷	32 ¹⁷
<i>Streptococcus pyogenes</i>	0,5 ³	2 ¹⁶
<i>Neisseria gonorrhoeae</i>		0,1 ¹⁶
<i>Haemophilus influenzae</i>		16 ¹⁶
<i>Escherichia coli</i>	0,12 ¹⁷	0,25 ¹⁷
<i>Enterobacter cloacae</i>	0,25 ¹⁷	2 ¹⁷
<i>Proteus mirabilis</i>	1 ¹⁷	2 ¹⁷
<i>Proteus constans (Providencia)</i>	1 ¹⁷	> 128 ¹⁷
<i>Salmonella typhi</i>	0,8 ¹⁵	1 ¹⁶
<i>Shigella dysenteriae</i>	0,8 ¹⁵	0,8 ¹⁶
<i>Yersinia enterocolitica</i>		< 2 ⁶

ESBL: Optionen orale Therapie ?

■ Orale Therapie (UTI)

- | | | |
|------------------------------|-----|---|
| ■ ? Fosfomycin <i>po</i> | 95% | } |
| ■ ? Nitrofurantoin <i>po</i> | 90% | |
| ■ ? Nitroxolin <i>po</i> | 90% | |
| ■ ? Pivmecillinam <i>po</i> | 85% | |
| ■ ? Cefpodoxim+(Amoxi-)Clav | | |
- in vitro-Empfindlichkeit



35.1
1 x 10 Tablets

Rx Cefpodoxime and Potassium Clavulanate Tablets

Microcef™ - CV 200

माइक्रोसेफ-सि वि २००

MICRO

Each film-coated tablet contains:
Cefpodoxime Proxetil IP
equivalent to
Cefpodoxime 200 mg
Potassium Clavulanate Diluted IP
equivalent to
Clavulanic Acid 125 mg

Colour : Titanium Dioxide IP

Dosage : As directed by the Physician.

Keep in a dry place, below 25°C

"SCHEDULE H DRUG-WARNING: To
be sold by retail on the Prescription of a
Registered Medical Practitioner only."

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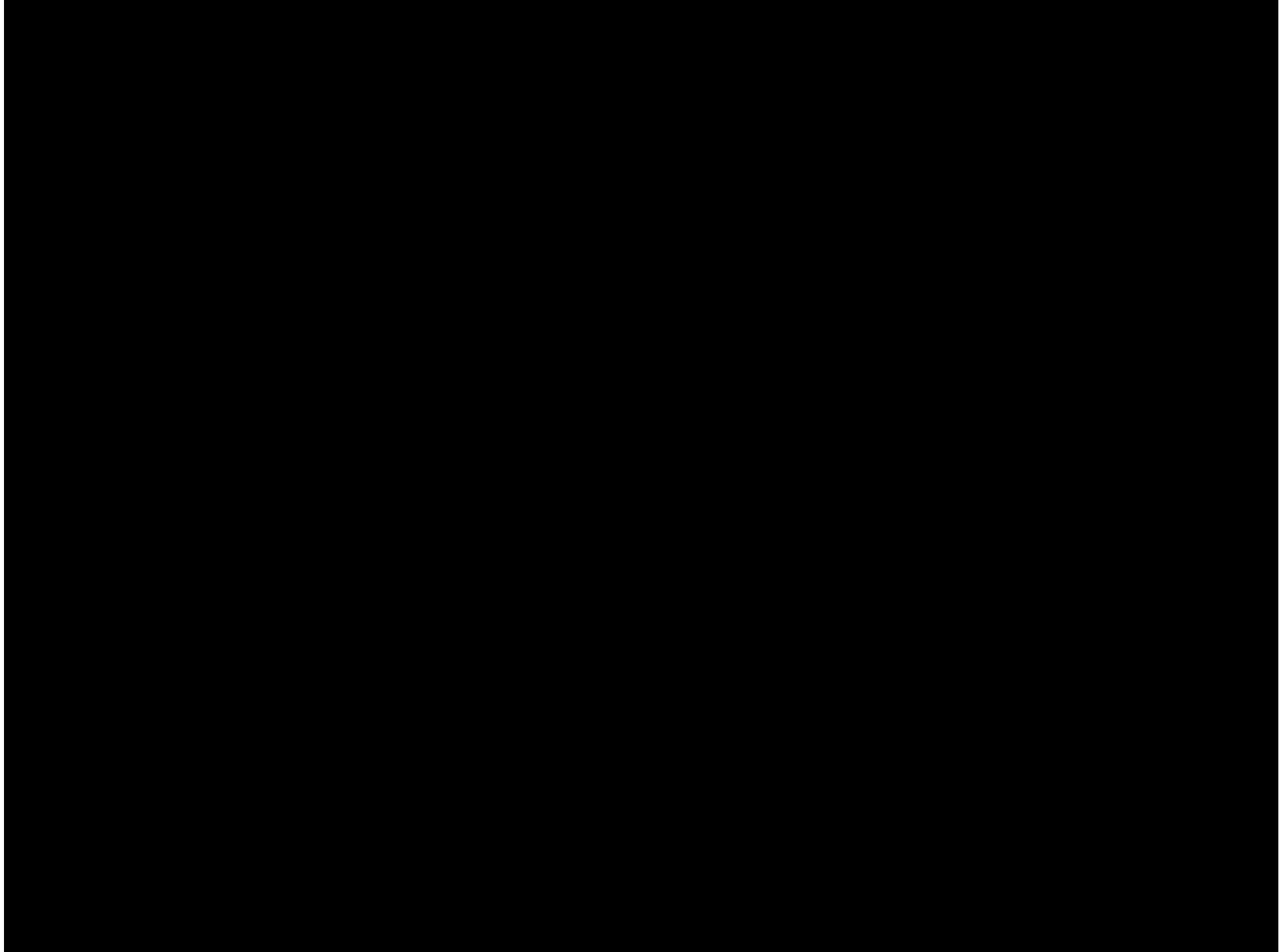
Batch No. : MVTB 0047

Mfg. Date : JUL . 2012

Expiry Date : DEC . 2013

Maximum Retail Price
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Inclusive of all taxes.
218 . 00

ESBL+ *Enterobacteriaceae*: Dekolonisierung ??



danke für's Zuhören