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IDs and CMs working together and in multidisciplinary teams

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Fondo Europeo de Desarrollo Regional
"Una manera de hacer Europa"



Why do IDs and CMs work together?

- Timely initiation of effective antimicrobial therapy is crucial to improving patient prognosis
- Antibiotic treatment should be accelerated, ceased or de-escalated according to clinical outcome and microbiological results
- Rapid detection of MR transmission is crucial
- Good clinical information permits low-cost microbiological diagnosis

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Case 1

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Case 1

- ❖ June 15th, 2016: Daily IDs and CMs meeting (11.30 am) → ***Streptococcus* spp. in BC (2/4 bottles)**
 - ❖ General Surgery ward
 - ❖ Male, 70 y. Colorectal carcinoma (2016). Surgery in May, 2016
 - ❖ Anastomosis dehiscence → tertiary peritonitis → 2nd surgery → ICU
 - ❖ Treated with piperacillin/tazobactam, and later meropenem for 16 days.
 - ❖ Patient admitted to surgery ward from ICU without antibiotics.
 - ❖ June 14th → 38.5°C, shivering → blood cultures → meropenem 1g/8h.
 - ❖ **Previous** cultures (abscess): *K. pneumoniae*, ESBL-*E. coli*.



2nd surgery



12 h

14th



38,5°C
Shivering



Bloodculture+**Meropenem**

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Case 1

- ❖ 12.10 am → ID to surgery ward:
 - ❖ 38.2°C in this moment. Sweaty. Blood pressure: 96/57 mmHg. 130 bpm.
 - ❖ Not feeling well. Pain under the surgical wound.

- ❖ **Taking into account the clinical story and present clinical situation....**
 1. Wait until final identification
 2. Add vancomycin or teicoplanin
 3. Add daptomycin
 4. Nothing, *Streptococcus* spp. represents a colonization/contamination
 5. All of them

ICU

Surgical ward

2nd surgery

10 h

16 h

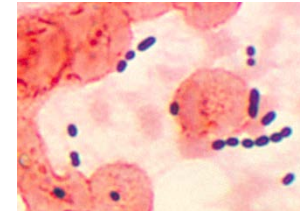
14th

PTZ

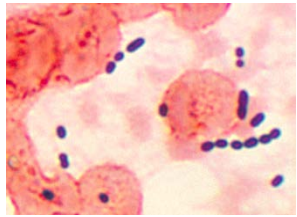
Meropenem

Blood culture

Positive



Gram positive cocci
Streptococcus



	Meropenem	Vancomycin
<i>S. pyogenes</i>	+++	+++
<i>S. agalactiae</i>	+++	+++
<i>S. mitis/sanguis</i>	+++	+++
<i>S. pneumoniae</i>	++?	+++
<i>E. faecalis</i>	+++	+++
<i>E. faecium</i>	+-	+++
<i>Peptostreptococcus sp</i>	+++	+++

ICU

Surgical ward

2nd surgery



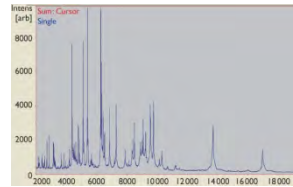
PTZ Meropenem

Bloodculture

GPC

Streptococcus

Maldi TOF



E. faecium

Blood:

E. faecium
Ampicillin R
Vancomycin S



Abdominal abscess → previous
ESBL *E. coli* + *E. faecium*

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Case 2

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Case 2

Hematological Ward

39°C, 96/55 mmHg, 123 bpm, sweating
PCT 17, lactate 3.5

1 d Neutropenia
(<500)

7 d

11:00

13:00

16:00

Patient 1

Acute leukemia

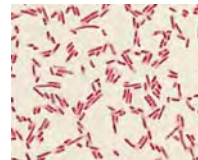
Admission for chemotherapy

Blood culture

Positive

Gram negative rods

Maldi TOF: *K. pneumoniae*



Case 2

❖ In this moment, you would...:

1. Start piperacilline/tazobactam 4g/8h
2. Start piperacilline/tazobactam 4g/8h + amikacin 1g/24h
3. Start ceftriaxone 1g/24h + metronidazole 500 mg/8h
4. Start meropenem 1g/8h
5. Start linezolid 600 mg/12 h
6. Wait and see...

Case 2

Antibiotic	MIC (mg/l)	Category
Amox/clav	>16/8	R
Piper/tazo	32	I
Cefuroxime	>16	R
Ceftazidime	>16	R
Cefotaxime	>16	R
Aztreonam	32	R
Cefepime	16	R
Ertapenem	>1	R
Imipenem	2	S
Meropenem	S	S
Gentamycin	>4	R
Tobramycin	>4	R
Trimetropin-sul	>4	R
Fosfomycin	>64	R
Colistin	≤ 0.5	S

Patient 1

Bacteremia
K. pneumoniae

- How often do you find this phenotype?
- Do you think there are some anomalous or unlikely results?
- Would you change the empiric therapy?

Case 2

Hematological Ward

Patient 2



9d

Patient 1

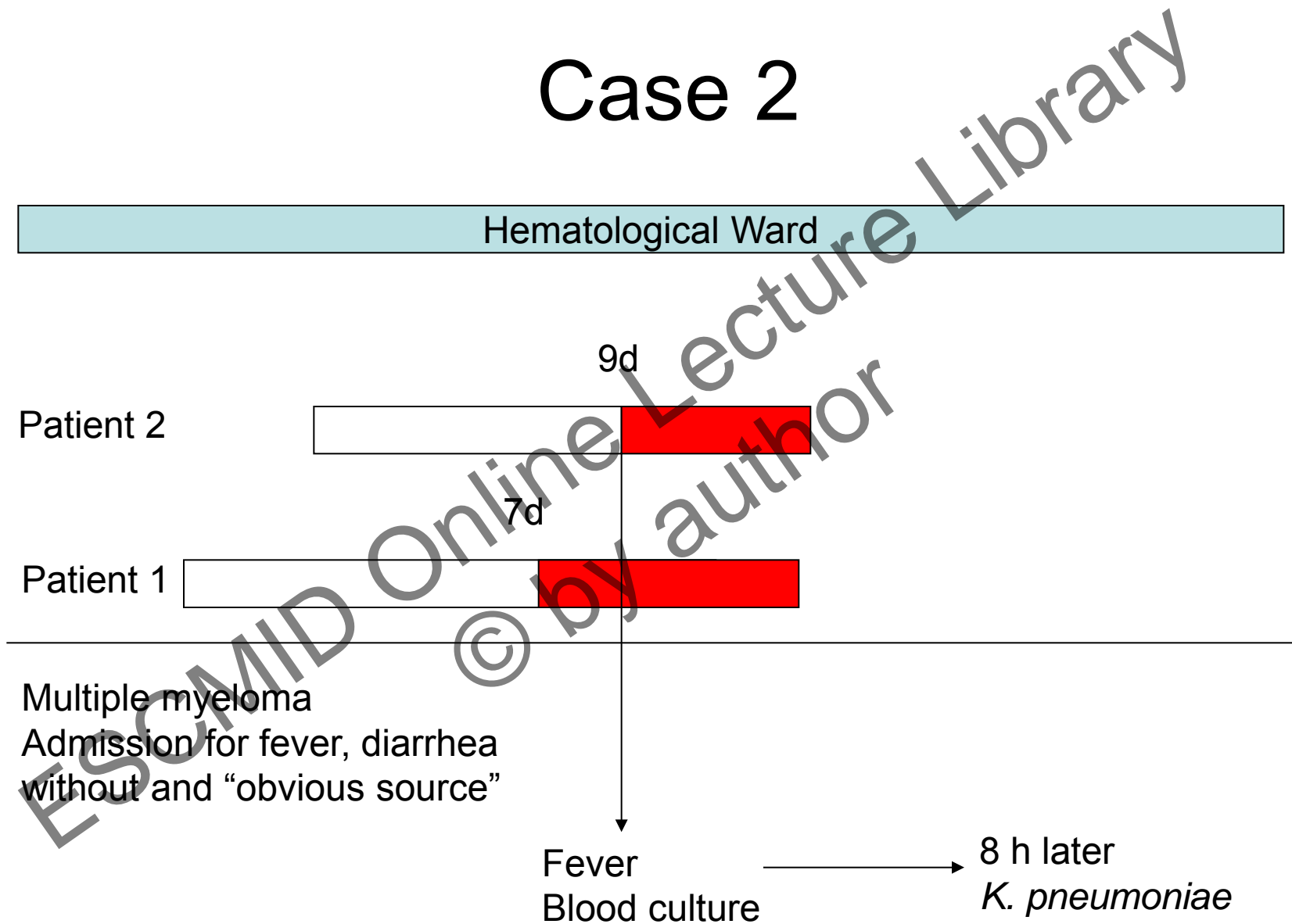


7d

Multiple myeloma
Admission for fever, diarrhea
without and "obvious source"

Fever
Blood culture

8 h later
K. pneumoniae



Patient 1

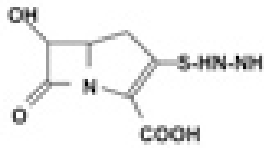
Patient 2

Antibiotic	MIC (mg/l)	Category	Antibiotic	MIC (mg/l)	Category
Amox/clav	>16/8	R	Amox/clav	>16/8	R
Piper/tazo	32	I	Piper/tazo	>64	R
Cefuroxime	>16	R	Cefuroxime	>16	R
Ceftazidime	>16	R	Ceftazidime	>16	R
Cefotaxime	>16	R	Cefotaxime	>16	R
Aztreonam	32	R	Aztreonam	32	R
Cefepime	16	R	Cefepime	16	R
Ertapenem	>1	R	Ertapenem	>1	R
Imipenem	2	S	Imipenem	4	I
Meropenem	2	S	Meropenem	2	S
Gentamycin	>4	R	Gentamycin	>4	R
Tobramycin	>4	R	Tobramycin	>4	R
Trimetropin-sul	>4	R	Trimetropin-sul	>4	R
Fosfomycin	>64	R	Fosfomycin	>64	R
Colistin	<=0.5	S	Colistin	<=0.5	S

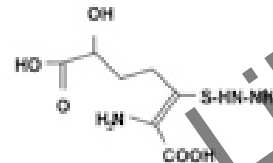
Some epidemiological questions

- ❖ Both isolates are very similar, so can we assume direct transfer between patients?
 - Outbreak?
 - Likely reservoirs?
 - Other measures?
 - Both isolates show resistance to ertapenem
 - Are they producers of carbapenemases?

Rapid microbiological tests



Hydrolysis of imipenem
+ pH indicator

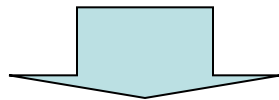


Color change

Name of test	Manufacturer	Time
Carba NP test	In house	2h 30 min
Blu Carba test	In house	2h
RAPIDEC® CARBA NP	BioMérieux	2 h
β CARBA™ test	Biorad	30 min

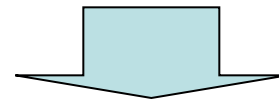


Isolate from patient 1



Negative

Isolate from patient 2



Positive

What would be your choice for targeted therapy in these cases?

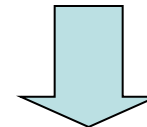
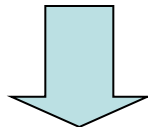
Patient 1

Patient 2

Antibiotic	MIC (mg/l)	Category	Antibiotic	MIC (mg/l)	Category
Piper/tazo	32	I	Piper/tazo	>64	R
Ertapenem	>1	R	Ertapenem	>1	R
Imipenem	2	S	Imipenem	4	I
Meropenem	2	S	Meropenem	S	S
Amikacin	1	S	Amikacin	1	S
Colistin	≤ 0.5	S	Colistin	≤ 0.5	S
Ciprofloxacin	>4	R	Ciprofloxacin	>4	R

ESBL+porin loss

ESBL+OXA-48



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Case 3

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Case 3

- ❖ June 7th, 2016: Daily IDs and CMs meeting (11.30 am)
 - ❖ Room 324/1: Gram negative bacilli in BC (3/4 bottles)
 - ❖ Male, 34 y. Diabetes, metastatic bladder carcinoma (2014) treated with chemo and radiotherapy. Charlson: 7
 - ❖ Double nephrostomy.
 - ❖ **Admitted on 6th**: Fever through 38.5°C, and obstruction of left nephrostomy.
 - ❖ No signs of severe sepsis. Leukocytosis, RCP 234 mg/L, normal renal function.
 - ❖ ER: ceftazidime 2g/8h was started.
- ❖ **Would you like to have any other information?**

Information about...

- a) Antibiotics in the last year?
- b) Last change of nephrostomy?
- c) Previous cultures?
- d) Known colonization?
- e) Previous admissions in the last year?
- f) Trips out of Spain?

Information about...

- a) Antibiotics in the last year?: ciprofloxacin (x2), ceftriaxone, fosfomycin and amoxicillin/clavulanate.
- b) Last change of nephrostomy?: 2 months ago.
- c) Previous cultures?: *E. coli* (salvage phenotype)
- d) Known colonization?: no.
- e) Previous admissions in the last year?: 4 admissions.
- f) Trips out of Spain? No.

Case 3

Male, 34 y.

Diabetes, metastatic bladder carcinoma (2014) treated with chemo and radiotherapy. Charlson: 7. Double nephrostomy

Day 0

Day 1

CEZTAZIDIME

Blood culture



Fever 38.5°C,

Gram negative bacilli

obstruction of left nephrostomy.

No signs of severe sepsis. Leukocytosis, RCP 234 mg/L, normal renal function

Case 3

- ❖ Fell not bad. Sitting on the bed.
 - ❖ Hemodinamically stable. 105 bpm, 18 bpm.
 - ❖ Obstruction has been solved by urology team.
- ❖ **Taking into account the clinical story and present clinical situation and new microbiological data....**
1. Would you change the empirical therapy?
 2. Wait until identification?
 3. Wait until susceptibility data?
 4. Add amikacin 15-30 mg/kg/day 1 dose?
 5. Change to carbapenems?

ESBL risk factors?

- a) Nephrostomy carrier
- b) Diabetes mellitus
- c) Previous admissions in the last year
- d) Charlson Comorbidity ≥ 4
- e) Previous therapies with betalactams or quinolones
- f) All of them

Specie-specific resistance determinants

- ❖ Do you know any characteristic of *S. marcescens*?...

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Case 4



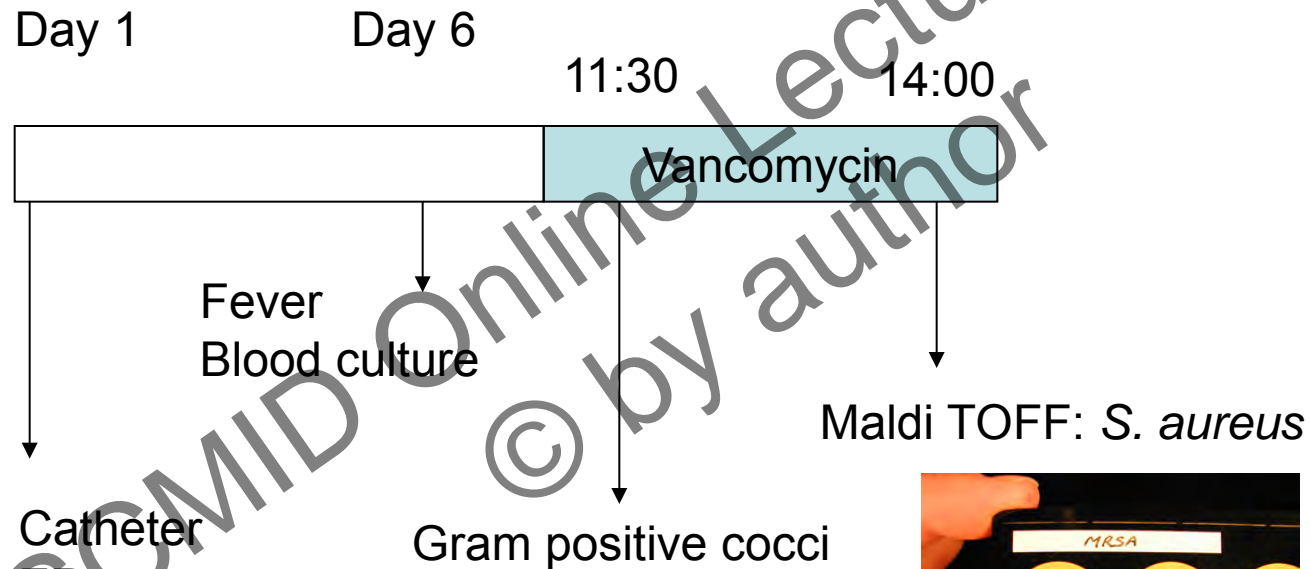
Case 4

- ❖ June 22nd, 2016: Daily IDs and CMs meeting (11.30 am)
 - ❖ Room 348/2: Gram positive coccus in BC (3/4 bottles)
 - ❖ Man, 48 y. Smoker, diabetes, hypertension, admitted because of an acute myocardial infarction
 - ❖ **Admitted on 16th**: peripheral catheter in ER without later change
 - ❖ Evaluated by Internal Medicine doctors the previous day ceftazidime 2g/8h + vancomycin 1g/12h was started
 - ❖ But...they did not remove the catheter because it hadn't "signs of phlebitis"
- ❖ June 22nd: 38.5°C, some shivering, BP 100/65 mmHg. No phlebitis.
- ❖ **What would you do in this moment?**

Case 4

- a) Change vancomycin for daptomycin (stop ceftazidime)
- b) Add cloxacillin 2g/4-6h to vancomycin (stop ceftazidime)
- c) Add cefazoline 2g/8h to vancomycin (stop ceftazidime)
- d) Stop ceftazidime
- e) The empirical therapy is OK. Nothing to change

Case 4



Rapid detection of *mecA*: negative

Case 4

❖ Options:

a) Stop vancomycin

b) Continue with combined therapy until final susceptibility inform

c) Change to daptomycin

d) Change to linezolid

Case 4

- ❖ **A negative result for *mecA* detection can predict the success of cloxaciline?**
 - a. Yes, because all the MRSA have *mecA* and the susceptibility of the agglutination is very high.
 - b. I don't know the susceptibility of these types of test
 - c. There is other resistance determinants different to *mecA*

Case 4

- ❖ June 23th, 2016
 - ❖ And what about targeted therapy?

HEMOCULTIVO/LIQUIDOS ESTERILES

SE AISLA:

(1) Staphylococcus aureus.

Antimicrobiano

S/I/R

CMI (ug/ml)

Penicilina	R	<=.12
Amoxicilina/clavulan	S	<=4/2
Oxacilina	S	<=.25
Gentamicina	S	<=1
Tobramicina	S	<=1
Amikacina	S	<=8
Levofloxacina	S	<=1
Eritromicina	S	<=.5
Clindamicina	S	<=.25
Tetraciclina	S	<=1
Rifampicina	S	<=.5
Tmp/Smx	S	<=2/38
Linezolid	S	<=1
Daptomicina	S	<=1
Teicoplanina	S	<=1
Vancomicina	S	1

Case 4

Anything more to do beyond antibiotic therapy?

- a) Remove the catheter
- b) Follow-up blood cultures in 48-72 hours
- c) Rule out septic thrombophlebitis
- d) Transthoracic echo if fever persists > 3 days or persistent bacteraemia appears
- e) Adjust duration to clinical complexity (from 2 to 6 weeks)
- f) All of them

Questions??



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