P13 Clinical microbiology IV – examination of wound and bloodstream infections

To study: Your own protocols (especially Special bacteriology)

Wound infections

Task 1: Specimens in wound infections

Try to fill in the following table:

Type of wound	Superficial	Deep wound with	Deep wound with not	Wound with pus,		
	wound	amount of pus sufficient	sufficient amount of	possibly containing		
		for being sent as a liquid	pus	anaerobic bacteria		
Sampling method						
When a specimen from a wound is send to the laboratory, it is very important to fill in the request form,						
especially to write 1)		and 2	2)			

Task 2: Imprint method for superficial wound examination (moulage method)

a) Imprint method – performing

Perform the imprint method in pairs. Place a sterile filtration paper on your mate's forearm (instead of a superficial wound). Let it 10 seconds here, then using tweezers, transport it carefully to a Petri dish with nutrient agar. After that, remove it and throw it away.

In practice, the filtration paper is not discarded, but sent together with the agar plate to the laboratory. In the laboratory the filtration paper is placed to two or three more media: agar with 10 % NaCl, chromogenic URI medium etc. After that, all media are cultivated overnight.

b) Imprint method – reading of results

Try to read the result of imprint method on URIchrom chromogenic medium using recounting scheme on your table and with the help of the key of colours of individual bacteria on the chromogenic medium. Attention! You have real results from real patients. Your result is not supposed to be the same as the result of your neighbour with another agar plate. Even the number of strains may be different.

The cultivation result of my imprint contained:

Likely species of bacterium	Quantity (approx. number of colonies per 25 cm ²)
1.	
(2.)	
(3.)	

Clue for preliminary diagnostics: Staphylococci – white on URI, growing also on NACL, white colonies on blood agar; Haemolytic streptococci – haemolytic colonies on blood agar, not growing on NACL, on URI not growing or (S. agalactiae) pale blue. Enterococci have grayish colonies on URI and small, but clearly blue colonies on URI. Enterobacteriaceae and G- non-fermenters – growing on Endo agar. Escherichia is pink on URI, Klebsiella is blue on URI, Proteus is yellow on URI, Pseudomonas is white or slightly green (because of its own pigmentation) on URI. All this is only preliminary, the algorithms from previous practicals are valid!

Task 3: Deeper wound swab result

In the case of a wound swab, there is no "common flora". That is the main difference between wound swab and e. g. swabs from respiratory ways: it is not necessary to search for a pathogen among the normal flora. On the other hand, we mostly use more culture media to detect all possible pathogens, even if they would be in a mix of them. Besides blood agar and Endo agar we usually use also blood agar with 10 % NaCl and blood agar with amikacin in order to search for streptococci and enterococci (but none of these media is used in our task). In other situations there is one pathogen only, and even in small amounts, so we have to multiply it in a liquid medium (broth). Also this medium is not present in our task.

Fill in the form again.

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Kód pojišťovny požadkuja IČ	7 2 1 2 3 4 5 6 Datum	Čís. dokladu	1, 2, 4, 4	
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POUKAZ NA VYŠE	TŘENÍ / OŠETŘENÍ	IČP		
Pacient Lucy Yellow	~~	Odbornost		STOR
Č. pojištěnce *1983	Dg: Suppurating wound of	Var. symbol		
Variabilní symbol	planta pedis	Datum	Kód	Poč
Odeslán ad:				
	Kód náhrady	2		
Požadováno:				
Nound with pus on	planta pedis, caused			1. 1.
by stepping on a tin i	n a pond;	5		
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ine pas appearea a	iter two days	7		18.00
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Patient: Lucy Yellow *1984 Dg.: wound of planta pedis							
Specimen: wound swab* Ordered by: Dr. Microbe Terrible							
*note: pyogene wound on planta pedis, swimming in a pond							
Growth on blood a. (incl. sme	ll) Endo agar:	MH agar:		Oxidase:	Conclusion:	Interpretation	
Antibiotic susceptibility test Final conclusion and for treatment:				and recommen	dment		

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Bloodstream infections

Task 4	Dlaad	a 14a	
1 ask 43	Biooa	cuitures –	processing

Describe the use	e of three types of blood culture vessels.
Fill in which of	data should not be missing on the order form in the case of blood culture (only "material
type/examination	on type" field)
Evoloin	
Explain: Why is absolute	e sterility in blood culture samples more necessary than in any other blood specimens (e. g. those
	nical examination)?
How many bloc	od cultures should be taken and why?
Tio Williamy 6166	a carrares should be unterfully willy.
Fill in the missi	ing fields in the description of blood culture processing and examination according to the video
clin and the tead	Pher's explanation
A blood culture	evessel arrives in the laboratory. Here it is put into a
The positive re	esult is demonstrated by and
When the cultiv	vation is positive, a smear is prepared and the content of the vessel is
onto the blood a	and Endo agar. Also, a preliminary test is performed directly
from the specim	nen; as the inoculum is not standardized here, its results are only
•	,
Task 5: Bloo	od cultures – microscopy of a positive specimen
	for blood cultures revealed a positive result. For preliminary treatment, a Gram stained smear is
	the contain of the vessel. Observe the result and write it. Attention! The slides have origin in ures of different patients. Therefore your result may be simply different from that of your
	a different slide.
Dland gultura a	ontained gram-positive – gram-negative* cocci – bacilli* arranged in
	copriate **only for cocci (pairs, chains, clusters) or G+ bacilli in palisades
	od cultures – cultivation result
	ation result of a positive blood cultures inoculated on solid media. Suggest more methods for
	ostics of bacteria. Try to assess preliminary antibiotic susceptibility. Also here you are not
Name of mediu	ve the same results as your neighbour.
Growth Y/N, ap	
of colonies	
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More tests of more detailed determination:							
Preliminary name of the microbe:							
Susceptibility testing Name of the set of antibiotics:							
Antibiotic				Antibiotic			
	ce	þ	Susceptible – resistant		ce	þ	ible ınt
	ren	sure	ısceptibl resistant		ren	sure	ısceptibl resistant
	Reference size	Measured size	usc res		Reference size	Measured size	Susceptibl – resistant
	R Si	N Si	·		R	N Si	
1.			S - R	4.			S - R
2.			S – R	5.			S – R
۷.			5 – K	J.			5 – K
3.			S-R	6.			S - R

Task 7: Blood cultures – interpretationFind suitable interpretation for results of two different patients.

John White, *1942, elevated temperature and inflammatory markers, three blood culture specimens sent to the laboratory	Joe Black, *1945, elevated temperature and inflammatory markers, three blood culture specimens sent to the laboratory
I Central venous catether. Time to detection 10 hours,	I Central venous catether. Time to detection 8 hours,
finding: Staphylococcus hominis, susceptible to	finding: Staphylococcus epidermidis, susceptible to
oxacilin, tetracycline, vankomycin, resistant to	oxacilin, resistant to tetracycline, vankomycin,
erythromycin, klindamycin, co-trimoxazole.	erythromycin, klindamycin, co-trimoxazole.
II Peripherial catather. Time to detection 13 hours,	II Peripherial catather. Time to detection 26 hours,
finding: Staphylococcus hominis, susceptible to	finding: Staphylococcus hominis, susceptible to
oxacilin, tetracycline, vankomycin, resistant to	oxacilin, tetracycline, vankomycin, erythromycin,
erythromycin, clindamycin, co-trimoxazole.	clindamycin, co-trimoxazole, no resistance observed
III Venepunction. Time to detection 13.5 hours,	III Venepunction. Time to detection 38 hours, finding:
finding: Staphylococcus hominis, susceptible to	Staphylococcus epidermidis, susceptible to oxacilin,
oxacilin, tetracycline, vankomycin, resistant to	co-trimoxazole, vankomycin, resistant to tetracycline,
erythromycin, clindamycin, co-trimoxazole.	erythromycin, clindamycin.
Likely interpretation:	Likely interpretation:

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