PZ13 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 fr | om a wound | is send to the laboratory, i | t is very important to f | ill 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

A sterile filtration paper on is placed on a superficial wound. We let it for 10 seconds here, then using tweezers, we transport it carefully to a Petri dish with nutrient agar. After that, the filtration paper is 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. Dental students do not perform this part practically.

b) Imprint method – reading of results

Try to read the preliminary 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. More precise determination and antibiotic susceptibility test would not be performed in this task.

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 greyish colonies on blood agar 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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| (protestation) | 2 1 2 3 4 5 6 Datum | Čís. dokladu | |
|--|--------------------------|------------------|---------|
| 1 1 1 Odborn | | (d) B proved) | Poř. č. |
| POUKAZ NA VYŠETŘE | NÍ / OŠETŘENÍ | IČP | |
| Pacient Lucy Yellow | | Odbornost | |
| Č. pojištěnce *1983 | Dg: Suppurating wound of | Var. symbol | |
| Variabilní symbol | planta pedis | Datum | Kód Poč |
| Odeslán ad: | Kód náhrady | | |
| Požadováno: | - | 3 | |
| Wound with pus on pla by stepping on a tin in a | | 4 | |
| the pus appeared afte | | 6 | |
| Poznámka: | | 8 | |
| 72 Dr. Microbe Temble 123 general Aracut oner | One: | 10 | |
| 456 (Campositive 8, Brno | - | 12 | |
| VZP-06x/1999 | | 14 | |

| Patient:Luc | су Үе | ellow | *198 | 4 Dg.:w | ound of pla | nta pedis | |
|---|------------------------------------|-------|--------------------------------|---------|----------------------|----------------|--|
| Specimen: wound swab* Ordered by: Dr. Microbe Terrible | | | | | | | |
| *note: pyogene wound on planta pedis, swimming in a pond | | | | | | | |
| | | | | | | Interpretation | |
| Antibiotic susceptibility t Piperacillin+tazobactam (TZP) | $C \ge 18$ $R < 18$ | | Ciproflox (CIP) | acin | C ≥ 25 R < 22 | | |
| Gentamicin (CN) Ofloxacin | $C \ge 15$ R < 15 $C \ge 16$ | | Ceftazidi (CAZ) Colistin | me | C≥16 R<16 C≥11 | | |
| (OEL) | D < 13 | | (CT) | | D < 11 | | |

(OFL)R < 13(CT)R < 11write S = susceptible, R = resistant, eventually I = intermediary

Final conclusion and recommendation for treatment:

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^{*}result of this test is also valid for doxycycline

Bloodstream infections

| Task 4: | Blood | cultures - | processing |
|----------|-------|-------------|--------------|
| I ask T. | Diou | cuitui cs — | DI OCCSSIIIZ |

| Describe the use | of three t | ypes of blood culture vessel | S. | |
|-------------------------------------|------------|------------------------------|--|---|
| | | | | |
| | | | | |
| | | | | |
| | | | | |
| Fill in which d type/examination | | | order form in the case of b | plood culture (only "material |
| J.1 | 71 | | | |
| | | | | |
| | | | | |
| Explain: | | | | |
| | | | re necessary than in any other | blood specimens (e. g. those |
| sent for biochem | nical exam | anation)? | | |
| | | | | |
| | | | | |
| | | | | |
| How many blood | d cultures | should be taken and why? | | |
| 110 11 1114111 9 1100 | | one and ever union and wing. | | |
| | | | | |
| | | | | |
| | | | | |
| | | | | |
| | | | culture processing and exami | ination according to the video |
| clip and the teac | ner's expl | anation. | it is put into a | |
| | | | | |
| The positive re | sult is de | emonstrated by | and | . |
| When the cultiv | ation is p | ositive, a smear is prepared | and the content of the vess | el is |
| onto the blood a | nd Endo a | ngar Also a preliminary | | test is performed directly |
| | | | | test is performed directly |
| from the specim | en; as the | inoculum is not standardized | d here, its results are only | · |
| | | | | |
| | | res – microscopy of a p | | |
| | | | | ment, a Gram stained smear is ion! The slides have origin in |
| | | | | different from that of your |
| neighbour with a | | | , | |
| D1 1 1/ | | •,• | ψ ' 1 '11'ψ 1' | in ** |
| | | | re* cocci – bacilli* arranged i s, clusters) or G+ bacilli in | |
| | _ | - | | pansaucs |
| | | res – cultivation result | | a Suggest more methods for |
| | | | | a. Suggest more methods for bility. Also here you are not |
| | | e results as your neighbour. | antibiotic susception | inty. This here you are not |
| Name of mediur | n | , , | | |
| Growth Y/N, ap | pearance | | | |
| of colonies | | | | |
| | | | | |

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| More tests of more detailed determined the second s | minatio | n: | | | | | |
|--|----------------------------------|---------------|-------------------|------------|----------------------------------|---------------|-------------------|
| Preliminary name of the microbe: Preliminary antibiotic susceptibe Name of the set of antibiotics: | | sting | | | | | |
| Antibiotic | Susceptibility Interpretation | Measured size | Result (encircle) | Antibiotic | Susceptibility Interpretation | Measured size | Result (encircle) |
| 1. | R < S ≥ | | S–I–R | 4. | R < S ≥ | | S-I-R |
| 2. | R < S ≥ | | S–I–R | 5. | R < S ≥ | | S–I–R |
| 3. | R < | | S–I–R | 6. | R < | | S-I-R |

Task 7: Blood cultures – interpretation
Look at 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: |
| Probably bacteriaemia | Probably pseudobacteriaemia |
| | |

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