

FACULTY OF DENTISTRY
ENGLISH LANGUAGE PROGRAM
 2020/2021. Fall semester
MICROBIOLOGY LABORATORY PRACTICES

1.	08, 09. Sept.	<p>Introduction to basic microbiology, laboratory rules.</p> <p>Microscopic examination of microbes</p> <ul style="list-style-type: none"> * Wet mount, vital staining of yeast * Preparation of smears (<i>S. epidermidis</i>, <i>E. coli</i>, <i>B. cereus</i>) * Simple and Gram staining * Preparation of smears from the surfaces of the tooth
2.	15, 16. Sept.	<p>Cultivation of bacteria</p> <ul style="list-style-type: none"> * Culture media (broth, agar-, blood agar- and chocolate agar plates, slant and stab forms) samples. Differential and selective culture media * Cultivation of anaerobic bacteria (supplemented thioglycolate and chopped meat medium, GasPack jar), Anaerobic chamber * Colony morphology of bacteria (<i>S. epidermidis</i>, <i>S. aureus</i>, <i>S. pyogenes</i>, <i>S. mitis</i>, <i>E. coli</i>, <i>Klebsiella</i>, <i>P. aeruginosa</i>, <i>Proteus</i>, <i>Serratia</i>, <i>B. cereus</i>, <i>H. influenzae</i>) * Procedure for making single colonies on plates <i>S. epidermidis</i> * Transport media, bloodculture (haemoculture)
3.	22, 23. Sept.	<p>Methods for sterilization and disinfection</p> <p>Macroscopic evaluation of the cultures of the previous practice</p> <ul style="list-style-type: none"> * Control of sterility (culture media: thioglycolate and glucose containing broth media) * Examination of the effect of an antiseptic on the bacterial flora of the skin (culturing before and after hand washing in the antiseptic solution on agar plates) * Microbial corrosion
4.	29, 30 Sept.	<p>Macroscopic evaluation of the cultures of the previous practice</p> <p>Antibiotic susceptibility of microbes</p> <ul style="list-style-type: none"> * Measurement of antibacterial activity (tube dilution method, plate dilution tests. Diffusion tests: well, and filter paper disc) * Examples for antibiotic susceptibility (<i>S. aureus</i>, <i>E. coli</i>, <i>S. pyogenes</i>, <i>P. aeruginosa</i>) * E -test, kits * Determination of antibiotic susceptibility of different bacteria (<i>S. aureus</i>, <i>E. coli</i>, <i>P. aeruginosa</i>) * Determinations of MIC and MBC for bacteria * Monitoring test of the antibiotic level in the serum

5.	06, 07. Oct.	<p>In vitro antigen-antibody reactions (serological methods)</p> <ul style="list-style-type: none"> * Macroscopic evaluation of the cultures of the previous practice Agglutination (qualitative: serotyping of <i>E. coli</i> on slide, quantitative: tube dilution test) * Precipitation (ring test, agar gel diffusion test, immun-electrophoresis) * Indirect agglutination * Complement fixation (CF): titration of complement * Enzyme-linked immunosorbent assay (ELISA) * Immunofluorescent methods
6.	13, 14. Oct.	<p>Midterm I. (General bacteriology and immunology)</p> <p>PYOGENIC COCCI</p> <p>Gram-positive cocci</p> <ul style="list-style-type: none"> * Cultures: <ul style="list-style-type: none"> ○ agar plates: <i>S. epidermidis</i>, <i>S. aureus</i> ○ blood agar plates: <i>S. epidermidis</i>, <i>S. aureus</i>, <i>S. pyogenes</i>, <i>S. pneumoniae</i>, <i>S. mitis</i>, <i>E. faecalis</i> ○ chocolate agar plate: <i>S. pyogenes</i>, <i>S. mitis</i>, <i>S. pneumoniae</i>, <i>E. faecalis</i> * Smears: <i>S. aureus</i>, <i>S. pyogenes</i>, <i>S. pneumoniae</i> (Gram stained) * Enzyme reactions: <ul style="list-style-type: none"> ○ catalase test on slide (<i>S. aureus</i>) ○ DNase production (<i>S. aureus</i>) ○ Coagulase test (<i>S. aureus</i>) * <i>S. mutans</i> on MS medium * Antibiotic susceptibility of <i>S. aureus</i>, <i>S. pyogenes</i> <p>Gram-negative cocci</p> <ul style="list-style-type: none"> * Culture: <i>N. pharyngitidis</i> on agar plate * Smears: Gram-stained smear of <i>Neisseriae</i> and methylene-blue stained smear of gonorrhoeal exudate * Enzyme reaction: oxidase test (<i>N. pharyngitidis</i>)
7.	20, 21. Oct	<p>Enteric Gram-negative rods (<i>Enterobacteriaceae</i>)</p> <ul style="list-style-type: none"> * Smears: <i>E. coli</i> (Gram stain), <i>Klebsiella</i> (capsule staining) * Cultures: agar plates: <i>E. coli</i>, <i>Klebsiella</i>, <i>Proteus</i> eosin methylene blue (EMB): <i>E. coli</i>, <i>Klebsiella</i>, <i>Proteus</i> * Selective cultures: deoxycholate-citrate (DC) agar: <i>Shigella</i> and <i>E. coli</i>, bismuth sulfite medium: <i>S. typhi</i> * Biochemical reactions: <ul style="list-style-type: none"> ○ carbohydrate fermentation in TSI media (<i>E. coli</i>, <i>Proteus</i>, <i>Klebsiella</i>, <i>Shigella</i>, <i>S. typhi</i> and <i>S. paratyphi</i>) ○ H₂S-, indol production ○ Urea degradation test * Serology: Widal test (tube dilution test: <i>S. typhi</i>) <p>Vibrios (sterile TCBS culture medium)</p> <p>Campylobacter spp., Helicobacter pylori (slides)</p> <ul style="list-style-type: none"> * Sampling containers <p>Gram-negative non-fermenting rods</p>

		* Pseudomonas on agar plate
8.	27, 28. Oct	<p>Gram-negative coccobacilli</p> <ul style="list-style-type: none"> * Haemophilus on chocolate agar plate and satellite phenomenon * Bordetella cultivation (sterile Bordet-Gengou medium) * Legionella (sterile BCYE medium) * <i>Actinobacillus actinomycetemcomitans</i> (smear from dental sulcus) Slides. <p>Mycobacteria</p> <ul style="list-style-type: none"> * Cultures :<i>M. tuberculosis</i> and <i>M. bovis</i>, atypical and apathogenic mycobacteria (Loewenstein-Jensen, Sula, Sauton and Dubos media) * Stained smear of sputum (acid-fast-staining) * Sampling containers
9.	03, 04. Nov.	<p>Gram-positive bacilli</p> <ul style="list-style-type: none"> * Cultures: <i>C. diphtheriae</i> on Löffler's coagulated serum slant medium and on tellurite plate * Smear of <i>C. diphtheriae</i> (stained) * Virulence tests: in vitro: agar gel precipitation <p>Endospore-forming Gram-positive aerobic and anaerobic bacilli</p> <ul style="list-style-type: none"> * Cultures: <i>B. cereus</i> on agar plate, <i>C. perfringens</i> and <i>C. tetani</i> in liquid media (containing thioglycolate or chopped meat) * Smears: Gram-stained preparations of endospore-forming bacteria <p>Gram-negative anaerobic rods</p> <ul style="list-style-type: none"> * <i>Bacteroides</i> * <i>Fusobacterium</i> * <i>Prevotella, Porphyromonas</i> <p>Gram-positive anaerobic rods</p> <ul style="list-style-type: none"> * <i>Lactobacillus</i> * <i>Bifidobacterium</i> * <i>Propionibacterium</i> * <i>Actinomyces</i>
10.	10, 11. Nov.	<p>Spirochaetales</p> <ul style="list-style-type: none"> * <i>Treponema pallidum</i> morphology and disease, Serology: non-treponemal antibody tests (flocculation test: VDRL, CF test) * <i>Borrelia</i> (slides) * <i>Leptospira</i> in Korthof medium and stained smear of <i>Leptospira</i> <p>Rickettsia</p> <ul style="list-style-type: none"> * Serology: Weil-Felix reaction <p>Chlamydia</p> <p>Mycoplasma (BEA, BEG cultures)</p>
11.	17, 18. Nov.	<p>Midterm II. Systemic bacteriology</p> <p>Medically important fungi</p> <ul style="list-style-type: none"> * Cultures: <i>C. albicans</i>, Penicillium, Aspergillus, Mucor on Sabouraud's agar plate * Block preparation of different molds * Smear: <i>C. albicans</i> (methylene-blue)

		Throat and nose bacterial samples, inoculation of blood and chocolate agar
12.	24, 25. Nov.	Macroscopic evaluation of the cultures of throat and nose samples. General virology <ul style="list-style-type: none"> * Cultivation of viruses: tissue cultures * Morphology of viruses * Cell-virus interactions (CP on tissue culture, HA in microplate) * Serological tests: HI, CF, IF, Paul-Bunnel test * Tests for detection of nucleic acid (PCR, hybridization) * Vaccines * Bacteriophages (morphology and phage typing) * Diagnosis of viral infections
13.	01, 02. Nov.	Most important DNS and RNS viral diseases <ul style="list-style-type: none"> * Viral diseases in oral cavity
14.	08, 09. Dec.	Medical parasitology (the most important parasitic protozoa and helminths): macroscopic and microscopic preparations Sampling containers. Summary of microbiological laboratory techniques and diagnostic procedures