

## **NFECTIOUS PROCESS**

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Infectious PROCESS is an interaction between micro- and macro-organisms (under the impact of natural and social factors of the environment).

Infectious DISEASE is a clinically marked part of this process.

## Infectious diseases

There is an agent =>

 Contagious: can be transmitted to another macro-organism => possibility of an outbreak.

Cyclic course (timing).

## Infectious process.



#### **Biological basis of infectious process**

#### **Agent's factors:**

pathogenic power; portal of entry of infection; dose

#### **Host's factors:**

 genetically determined: non-specific and specific resistance (HLA)
 acquired: nutrition, intoxications, ecologic factors, behavior patterns, vaccination, treatment.

## Complications

 Specific: typical to the disease (perforation of ulcers of small intestine in typhoid fever patients)

 Non specific (sepsis of another origin due to prolonged presence of intravenous catheter). Symptoms and signs of infectious diseases

Fever Rash Lymphadenopathy Liver /spleen enlargement Respiratory syndrome Diarrhea Hepatitis Meningeal syndrome, etc



## Syndromes

 Congunctivitis, Tonsillitis, pharyngitis, stomatitis, … Pneumonia, bronchitis... Gastro-entero-colitis... Hepatitis... Kidney insufficiency (acute, chronic) Meningitis... DIC, etc

## Diagnosis

- Anamnesis, symptoms and signs => <u>syndromes.</u>
  - <u>Prove the syndrome</u>: biochemical tests, ECG, X-ray, USI, etc.
- Anamnesis, association of syndromes => suggestion of etiology.
   <u>Clinical etiologic diagnosis is always</u> <u>hypothetical</u> => how to check it?

## Etiologic diagnosis

To prove or to disapprove it: to find the supposed agent or to find its markers.
 Markers: Ag of the agent or <u>Ab to it</u>.

 Methods depend on the agent: bacteria, virus, rickettsia, clamydia, mycoplasma, protozoa, helminthes, fungi.

## Microscopy

#### Pluses:

- fast

 the main method for protozoa, helminthes, fungi.

Minuses: for bacterial infections in the most cases it is a tentative method.But sometimes can be very informative (N.meningitidis in CSF).

**Bacteriological investigation** Pluses: accurate; sensitivity to antibiotics <u>Minus</u>: needs time (several days or more) Negative result does not always turn down a supposed diagnose: - defects of sample taking, transportation,

media and lab technique;

 recovery stage (spontaneous or due to correct treatment).

<u>Absence of correct suggestion! => media</u>

## Serological investigations

To detect antibodies to a <u>suggested agent</u>
Antibodies – <u>in serum</u> (CSF).

- <u>Pluses</u>: simple; reliable; cheap; often the only confirmation of a diagnosis. <u>Minuses</u>:
- "window period";
- investigation itself is fast, but results are always retrospective.

#### Antibodies



#### Antibodies



#### To prove etiological diagnosis: Ab

- 4 times increase in titers of Ab to the agent (primary or secondary immune response):
   Samples should be taken twice in time!
  - <u>1-st time: the 1-st week (zero is expected)</u>,
  - 2-nd time: in 2 weeks (maximum level).
- Diagnosis is late: after 2-3 weeks; can be even later under effective treatment =>
  - the 3d sample at week 5-6 of the disease.
- The only test can be (+) due to previous disease, vaccination, poly-agglutination.
   "Min diagnostic level of Ab" is not reliable.<sup>18</sup>

To prove etiological diagnosis: Ig
Ig M (+) to the agent even once means the primary immune response.
Ig M can be usually found since the 5-th day of the disease up to the 4-6 weeks.

Rare IgM can persist much longer (HBV).

 Ig G(+): >10 days of the disease (peak, recovery, chronic stage, previous disease or vaccination)-similar to Ab significance.

#### To prove etiological diagnosis: Ag

- Ag can be found in any substrate.
- No "window" period =>
- Express-techniques to reveal the Ag (Ab with some additional mark to make immune complex visible): plague, etc. PCR – to reveal DNA/RNA of the agent. <u>In blood PCR(+): replication; PCR(-):</u> no replication; sanitation -? => biopsy. Ag disappear in the process of sanitation in recovery stage => Ab.

#### Phases of the process

- The end of incubation and the first part of the disease – presence of Ag; no Ab: the most contagious and dangerous part. Recovery with clearing from the agent: all Ag disappear, Ab become (+). Chronic form: presence of Ag, or Ag+Ab; sometimes – only Ab (anti-HBcor Ab).
- Life prognosis depends mostly on tissues functions (biochemical tests, ECG, etc).

# Mixed infections, combination of different diseases

Confirmation of the one disease does not allow us to exclude another one.

 To exclude (or confirm) a disease we should investigate <u>for this disease</u>.



#### Exact diagnosis:

## Prognosis spontaneous course (subclinical, mild, moderate, severe), under the treatment

#### **Treatment**

- etiology,
- phase of the process,
- severity

## Infectious process.



#### Treatment

Etiotropic – to affect the agent.

 Pathogenetic (syndromic) – to improve or to replace tissues functions.

Symptomatic – to suppress symptoms.

#### **Etiotropic treatment**

Antibacterial, antiviral, antiprotozoal, etc.

Result of therapy depends mostly on - correct choice of spectrum and activity of preparations (if not correct: disease and treatment go own ways); when the treatment is started (the first 1-2 days => just stop the disease); duration of the treatment.

#### Pathogenetic (syndromic) treatment

- Can be life-saving (rehydration in cholera, hemodialysis in HFRS, dehydration in brain edema, intubation in laryngeal diphtheria).
- Often it is the main part of the treatment: DS is too late to start etiotropic treatment (HAV, HF), or etiotropic treatment is not correct, etc.

#### **Basic regiment**

 Bed rest
 Diet: in acute diseases – according to appetite; boiled and cultured milk foods can be used in any situation. Liquids.

 Clinical observation (behavior, t, pulse, BP, RR, diuresis, symptoms and signs).

Symptomatic treatment - can be useful.

