

Yersinia

Yersinias are pathogenic to humans. *Yersinia pestis* and *Yersinia pseudotuberculosis* have ecological and genetic similarities, but epidemiological differences. *Yersinia enterocolitica* is found mainly in water, soil and the digestive tract of vertebrates.

Yersinia pestis

Yersinia pestis is G- bacteria and a major pathogen causing **plague pandemics**. It was first isolated by Alexander Yersin in 1894 in Hong Kong. Plague is associated with high mortality without antibacterial therapy. In the wild, *Yersinia pestis* was found in several rodent species. However, *Yersinia pestis* circulates through flea bites, in humans it is a **rat flea** bite. Bacteria get into the wound. **Human-to-human** transmission can occur via **aerosol particles** or droplet infection and a primary plague infection arise. Plague is **rare** at this time, occurring in Africa, India, Southeast Asia, Mexico, and the western United States.^[1]

Morphology

Yersinia pestis is a pleomorphic **rod-shaped bacterium with a capsule**. It is a **stationary** stick. Other yersinias are mobile. A common feature of *Y. pseudotuberculosis* and *Y. enterocolitica* is that it **stains polar**. This rod, like all Enterobacteriaceae, is characterized by the ability to grow on simple laboratory soils. Their undemanding growth ranges from 0 to 40°C with a temperature optimum around 30°C.^[2]

Antigenic structure

On its surface is a localized **F1 protein complex**, which is a protective antigen. O-specific side chains are not present. Virulent strains form **V** and **W proteins**, the production of which is linked to plasmids.

Pathogenicity

The pathogenicity of *Yersinia pestis* is determined by a complex of chromosome and plasmid-linked factors. Surface factors that **block phagocytosis** are important. **Virulent strains** of this bacterium are **facultatively intracellular parasites** that multiply in macrophages. During infection, *Yersinia pestis*, an facultative intracellular bacterium, exhibits the ability to first invade cells and then thwart phagocytosis of the host cell. During these two distinct phases, the **invasion phase** and the **anti-phagocytic factor phase**, the bacteria in manipulating the host cell help to complete each of these functions, but the mechanism by which *Yersinia* regulates these functions during each step remains unclear. In addition to macrophages, the bacterium is also able to penetrate **epithelial cells**. During the invasion at the site of injury, a **hemorrhagic pustule** is formed, from where the infection is further distributed to the descending **lymph nodes**, in which the subsequent enlargement produces **smoke** (bubonic form of the plague). Later, **bacteremia** and sepsis develop. A secondary disease, Pneumonia, may develop, causing *Yersinia pestis* to spread to the surrounding area by coughing up, and the infected person will develop primary pneumonia – **a pulmonary form of the plague**. When a patient manages to overcome the disease, he develops good immunity.

Diagnosis and therapy

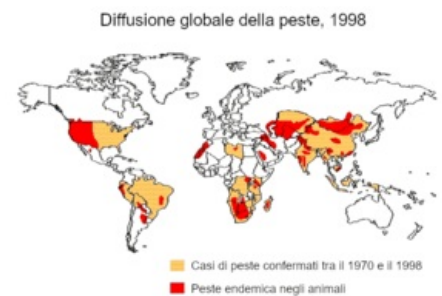
Bacteria are diagnosed from the contents of the pustules, lymph nodes, **blood culture**, bone marrow, and **sputum**. Cultivation is possible on blood agar. Bacteria stain according to Gram negative. When testing *Yersinia pestis* for antibiotic susceptibility, the bacteria were found to be **susceptible** to chloramphenicol, streptomycin, kanamycin, tetracycline, quinolones. However, plague vaccines produced by *Yersinia pestis* are in clinical development. Streptomycin can be used in endemic areas. We can induce **short-term immunity** when a **dead vaccine** with a protective antigen is given. A **live vaccine** with live attenuated strains is **more effective**, but the use of this vaccine is only approved in some parts of the world. There is currently **no licensed vaccine** to prevent plague in Western Europe.

Yersinia enterocolitica

Yersinia enterocolitica is a **G- non-sporulating, facultatively anaerobic** straight rod. It is pathogenic to humans and is the cause of human **yersiniosis**. The disease is transmitted alimentary through infected raw pork. It is widespread cosmopolitanly. The prevention is to avoid consuming badly roasted pork, or to observe hygiene after contact with the meat. The disease can be treated with antibiotics.



Yersinia pestis on blood agar



The spread of the plague in the world in 1998

Morphology

Yersinia enterocolitica is a **gram-negative straight** rod that is only **mobile** at lower temperatures. Its size is 0.3–1.0 µm and 1.0–6.0 µm. It can be grown on conventional cultivation soils because **it is easy to grow**. It has no problem growing in a low glucose environment at 28°C. It is recommended to use selective soils for culturing bacteria from faeces. The ideal pH for growth is in a wide range of 4.6-9 with an **optimal pH** of 7-8.

Epidemiology

Yersinia enterocolitica was discovered by Schleifstein and Coleman in year 1939 in the USA. It has been a well-known pathogen in the last four decades. It occurs mainly in food, especially in **raw pork**, but has no problem surviving on fruits and vegetables in the refrigerator at lower temperatures. Most *Y. enterocolitica* isolates recovered from natural samples, including houses where animals are slaughtered or in butchers. They were non-pathogenic in nature and in water. Bio serotypes 4/O:3 are the most common cause of human **yersiniosis**, food poisoning caused by *Y. enterocolitica* is known cosmopolitanly.



Yersinia enterocolitica

Pathogenicity and virulence

Virulence is bound on the chromosome and on the plasmid. It is given by the **invasiveness** and **penetration** of the microbe into the cell. Food can enter the terminal ileum and appendix, where it penetrates cells and lymphatic tissues to which it has a particularly high affinity. **Virulent** strains **multiply in macrophages** and induce **granuloma** formation. Non-virulent strains are eliminated without penetrating the cells.

Clinical picture

We observe clinical signs from **mild diarrhea** to serious complications such as **liver abscesses** and **post-infectious extraintestinal consequences**. The main reservoir is **pigs**, which are asymptomatic carriers.

Disease

Manifestations of the disease in humans are different in children and adults. Children develop **fever and diarrhea**, where blood may appear. **Abdominal pain** is also typical **in the right lumbar pit**. In adults, we observe **gastrointestinal infections** and diarrhea. With long-term illness, **secondary complications** can occur and these are **inflammations of the joints**. An unusual case of infection manifesting as **perianal ulcers** and colon ulcers has also been observed. An individual can become infected from infected pork. Toxic bacteria produce a **thermostable toxin** at 25°C in milk. However, at 4°C they will not produce significant amounts of this toxin.



Yersinia enterocolitica under a microscope

Therapy

Bacteremia requires **antibiotic** treatment. **Tetracycline** is used. Tests have shown that bacteria have a high degree of resistance to **ampicillin** and **cephalothin**. In the diagnostic laboratory, **examinations of the stool**, nodes, and appendix are performed. Immunochromatic test for the quantitative determination of serotypes O3 and O9 in a stool sample with a result within 1 hour. The test is highly sensitive and specific.

Prevention

Prevention against *Yersinia enterocolitica* and *Yersinia pseudotuberculosis* is to avoid eating undercooked pork or drinking unpasteurized milk. After contact with raw pork, it is recommended to wash your hands thoroughly with soap.

Yersinia pseudotuberculosis

Yersinia pseudotuberculosis is a G⁻, facultatively anaerobic, pleomorphic rod capable of moving at lower temperatures. It is widespread cosmopolitanly. Alimentary, orofecal transmission from infected animals or food. Diseases are associated with gastrointestinal problems, fever, diarrhea. The best prevention is personal hygiene and avoiding contact with infected animals.

Morphology

Yersinia pseudotuberculosis je **G–**, **pleomorphic rod**. It is able to **move** at lower temperatures. This organism was described in 1889 as a disease of **Guinea pigs**. However, *Yersinia pseudotuberculosis* has emerged as the ancestor of *Yersinia pestis*, which was the cause of pandemic plague during the years 541-767.

You can find more information on the page *Yersinia pestis*.

Epidemiology

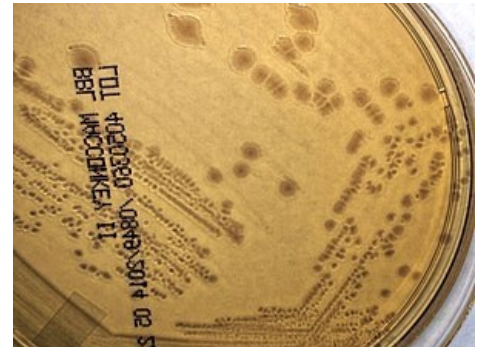
It is undemanding to cultivation, it is cultivated on common soils. It is widespread cosmopolitantly and has been isolated from several animal species such as cats, dogs, pigs, monkeys, and more. *Y. pseudotuberculosis* is widespread in the environment (feces, water), where it can survive for a long time. The environment itself is contaminated with the **faeces of infected animals**, especially rodents and birds. In addition to animals, it was also found in water, both in rivers and in alpine streams.

Antigenic structure

We can distinguish 5 serotypes according to somatic antigen.

Pathogenicity and virulence

The **toxin** bound in the cell is different from the plague toxin. Virulent strands can cause the **septicemic form akin to the plague** especially in individuals with a weakened immune system. Gastrointestinal tract diseases such as **acute** and **chronic appendicitis**, gastroenteritis, or disorders of mesenteric lymph nodes occur much more often. Infections with *Y. pseudotuberculosis* were reported all around the world. These infections happen less frequently than infections by *Yersinia enterocolitica*. Most of the infections are **sporadic** and arise rarely, e.g., outbreaks in Finland and Japan.



Yersinia pseudotuberculosis colonies on MacConkey-Agar (MAC) agar, showing a negative result for lactose fermentation.

Clinical manifestations

The disease manifests itself in humans with **fever**, rashes, **abdominal pain**, and **diarrhea**. *Yersinia pseudotuberculosis* has been associated with Kawasaki disease.

Diagnostics

Laboratory tests are obtained from **stool**, urine, or **blood tests** for bacterial antibodies. **Serotyping** of an isolated strain. Antibiotics are used for treatment. We must not forget that the patient may be dehydrated due to fever and diarrhea. **Rehydration** of the patient is necessary.

Links

Related articles

- Microbiology repetitorium

References

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2. BEDNÁŘ, Marek – SOUČEK, Andrej – FRAŇKOVÁ, Věra. *Lékařská mikrobiologie : Bakteriologie, virologie, parazitologie*. 1. edition. 1999. . ISBN 8023802976.

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- JULÁK, Jaroslav. *Úvod do lékařské bakteriologie*. 1. edition. Praha : Karolinum, 2006. ISBN 80-246-1270-4.