

# BETWEEN PIROPLASMOSIS AND ANAPLASMOSIS:

## A NUANCED DIAGNOSIS

### ISOLATED EQUINE FEVERS

**Equine vector-borne diseases**, such as **piroplasmosis** and **anaplasmosis**, are vector-borne-infections that are often responsible for similar clinical symptoms. However, these pathologies differ in their origin, geographical distribution and seasonality, making differential diagnosis essential to ensure rapid and effective treatment.

#### TWO DISEASES WITH SIMILAR SYMPTOMS BUT DISTINCT CAUSES

**Equine piroplasmosis** is caused by the **protozoan parasites** *Babesia caballi* and *Theileria equi*, transmitted by ticks of the genus *Dermacentor*, *Ixodes* and *Rhipicephalus*. When they bite, these ticks inoculate the parasites through their saliva, causing **clinical signs such as hyperthermia, despondency, anaemia and hepatorenal disorders in the horse**. In **acute forms**, the course can be **fatal in less than 72 hours** without appropriate treatment.

**Equine anaplasmosis**, on the other hand, is a **bacterial disease** caused by *Anaplasma phagocytophilum*, transmitted mainly by *Ixodes* ticks.

It causes **symptoms** similar to those of **piroplasmosis**, such as **fever, lethargy, anorexia, limb oedema**, and **thrombocytopenia**.

Often underdiagnosed, anaplasmosis is nonetheless present in Europe and North America, with distinct seasonal activity periods.

### GEOGRAPHICAL DISTRIBUTION AND SEASONALITY: A KEY FACTOR IN THE RISK OF INFECTION

**Piroplasmosis** is widely distributed in **tropical, subtropical and temperate regions**, including **South America, Africa, Asia** and **Southern Europe**. It tracks tick activity, with an increased **prevalence in spring and summer in temperate zones**.

**Anaplasmosis**, on the other hand, is more common in the **wooded and humid regions of Europe, North America, and Asia**. Its infection cycle is bimodal, with peaks in **spring and autumn**, when ticks of the genus *Ixodes* are most active.

### DIAGNOSTICS

During the **acute phase of infection**, tick-borne diseases can be diagnosed by various methods:



#### Serological method

Testing 2 weeks apart to check for seroconversion

- > By **indirect immunofluorescence**
- > By **fixing the complement**
- > By **ELISA techniques**  
(mainly used for chronic piroplasmosis)



#### Method of searching for the parasite (microscopy)

- > By blood smear (low sensitivity since the number of parasites is very low)



**Gene amplification** by PCR or LAMP is a great option. It offers **high sensitivity** as well as **simple and fast identification of the pathogen**.



**For maximum effectiveness**, it is recommended to combine several of these methods, as they are complementary and can detect different elements.



## TREATMENT

**Imidocarb** is a drug **that does not have an MA for equine treatment**. Its injection is very violent for the horse, leading to spasms, tremor, or convulsions.

**Digestive issues** are frequent, and colic or vomiting must be taken seriously as they can **result in fatal exhaustion**. Imidocarb should therefore only be used if a **positive case of piroplasmosis is established**.

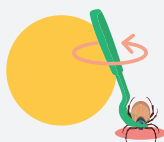
**Anaplasmosis** is often underdiagnosed and should not be treated with Imidocarb but with antibiotics. **Differentiating between piroplasmosis and anaplasmosis is essential for proper treatment.**

## HOW TO PREVENT ISOLATED EQUINE FEVERS?

**There is no vaccine against piroplasmosis or anaplasmosis in horses**, so here are some recommendations to give to your clients to avoid its occurrence:



**USE** an acaricide to be diluted in water and applied to the horses' backs to repel ticks. Avoid excessive use to prevent resistance.



**REMOVE** ticks promptly with a **tick remover** and monitor the bite site and the horse's overall condition over the following days.

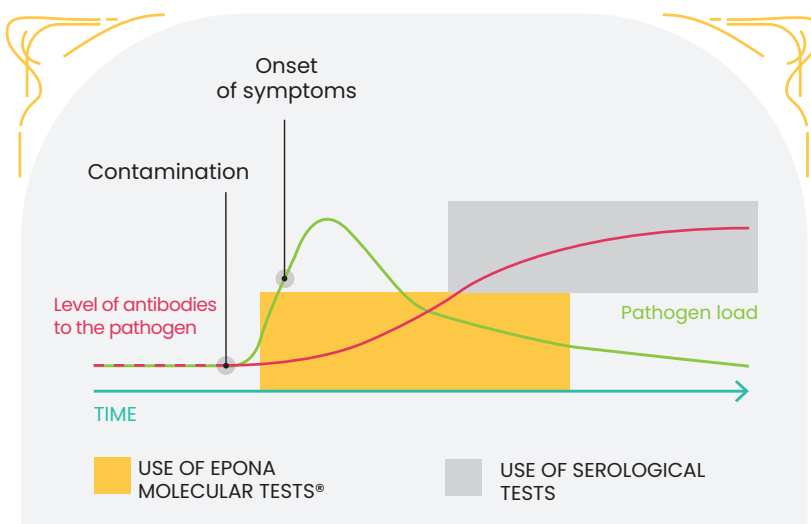


**MAINTAIN** their living space such as the hedges around the stud paddocks.

## WITH EPONA®: A FAST AND SENSITIVE DIAGNOSIS

Epona® tests are **molecular biology** tests that **directly detect the DNA of the pathogen** and can therefore be used from the **beginning of the infection** even before symptoms are detectable.

**Their sensitivity is very close to that of PCR**, but the results are obtained in **30 minutes**.



## COMPARISON OF THE DIFFERENT DIAGNOSTIC METHODS

	Microscopy in the clinic	Serology at the clinic	PCR in an external laboratory	Epona® Tests (LAMP-Technology)
Specificity	☐	☐	✓	✓
Accurate identification of the pathogen species	✗	✗	✓	✓
Sensitivity	☐	☐	✓	✓
Response time	✓	✓	☐	✓

