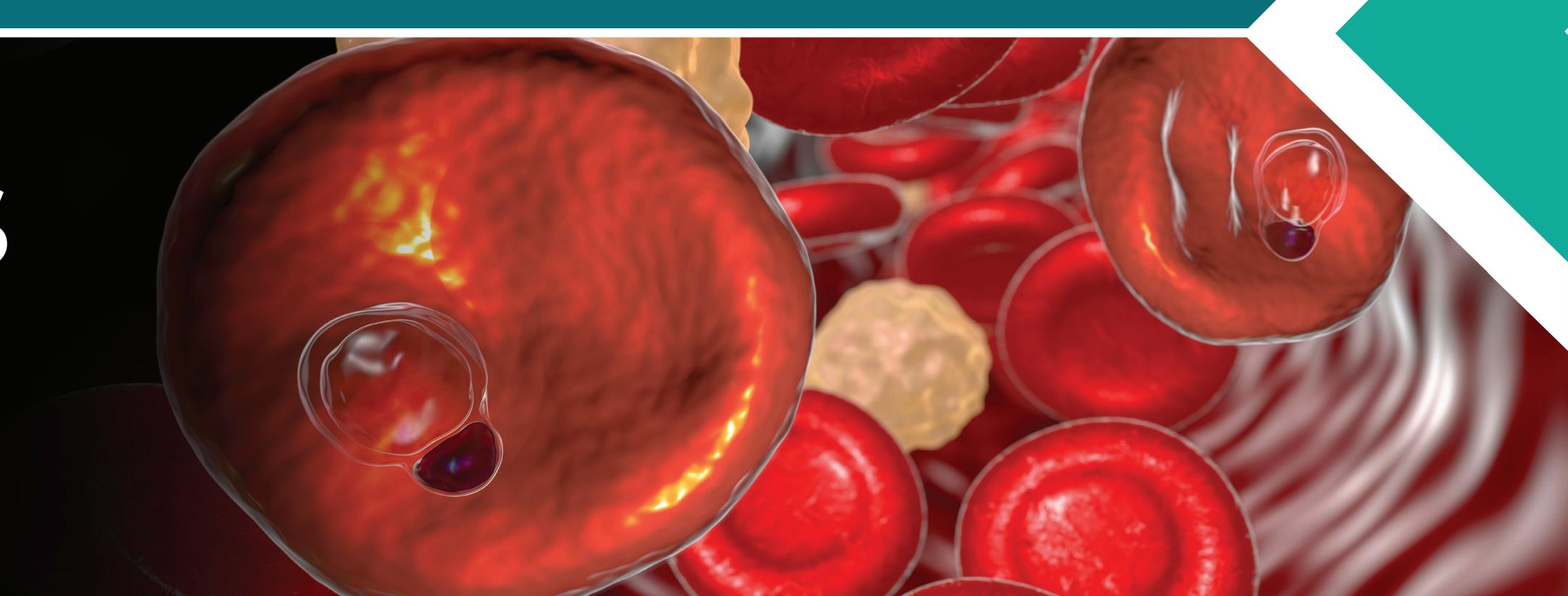


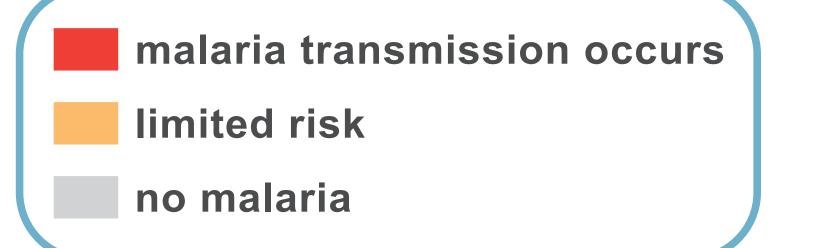
WHERE S MALARA FOUND?



GLOBALLY

The distribution of malaria depends mainly on climatic factors such as temperature, humidity, and rainfall.

Thus, in tropical and subtropical areas, where Anopheles mosquitoes can survive.



This map shows of the parts of the world where malaria transmission occurs.

LOCALLY

IN SOUTH AFRICA, MALARIA IS MAINLY TRANSMITTED ALONG THE BORDER AREAS

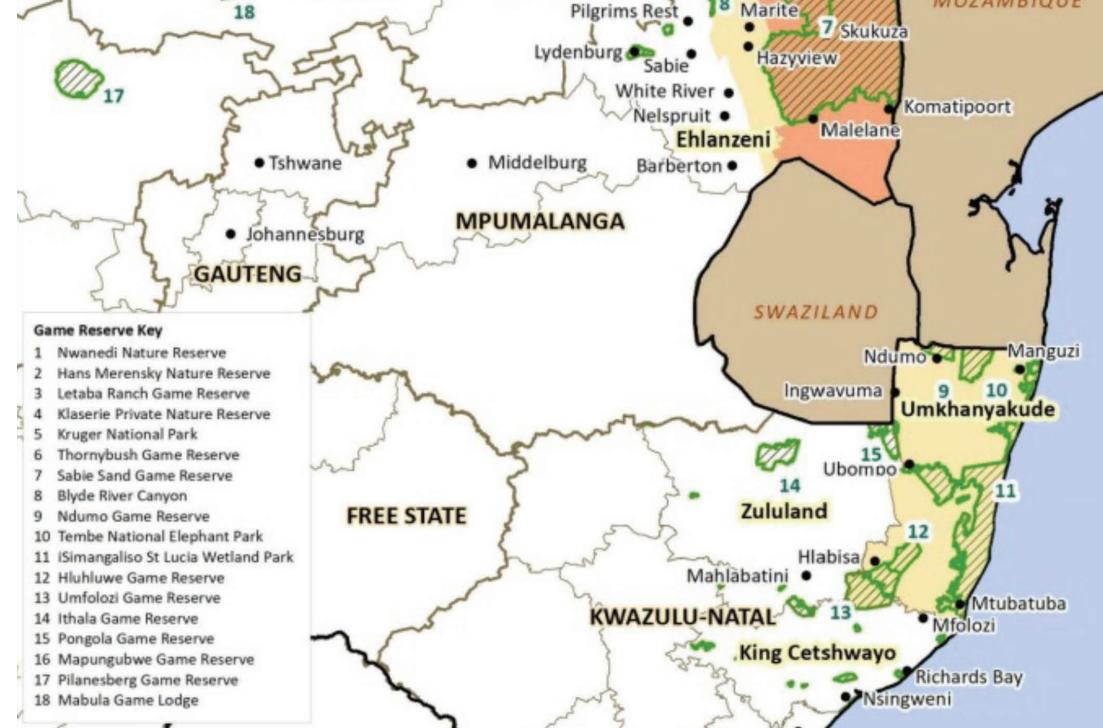
ZIMBABWE Alldays BOTSWANA Thohoyandou LIMPOPO ephalale 2 7 Capricorn Tzaneer Mopani Sekhukhur MOZAMBIQUE

10% of the population in South Africa (approximately **4.9 million** persons) is at risk of contracting malaria. Some parts of Limpopo, Mpumalanga and KwaZulu-Natal are endemic for malaria.

Low Risk

Only non-drug measures to prevent mosqito bites are recommended from September to May





Moderate Risk

Antimalarial drugs are recommended from September to May for all travellers

Malaria Risk does exist in neighbouring countries

For furthur information, please consult the WHO travel health guidelines.

Malaria transmission in South Africa is seasonal, with malaria cases starting to rise in October, peaking in January and February, and waning towards May.



HOW DOES MALARIA SPREAD?

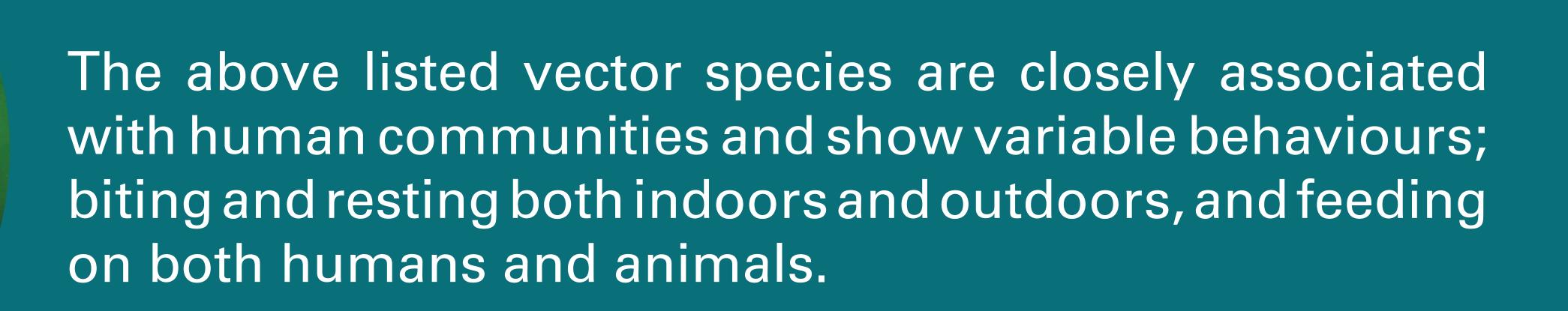
MOST COMMONLY BY THE BITE OF AN INFECTED FEMALE ANOPHELES MOSQUITO

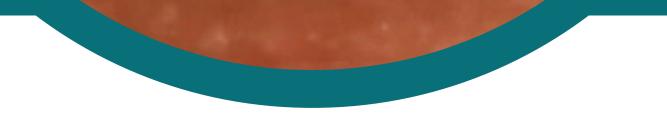
Only *Anopheles* mosquitoes can transmit malaria. However, not all *Anopheles* mosquitoes are malaria vectors, and not all vector species transmit malaria equally well.



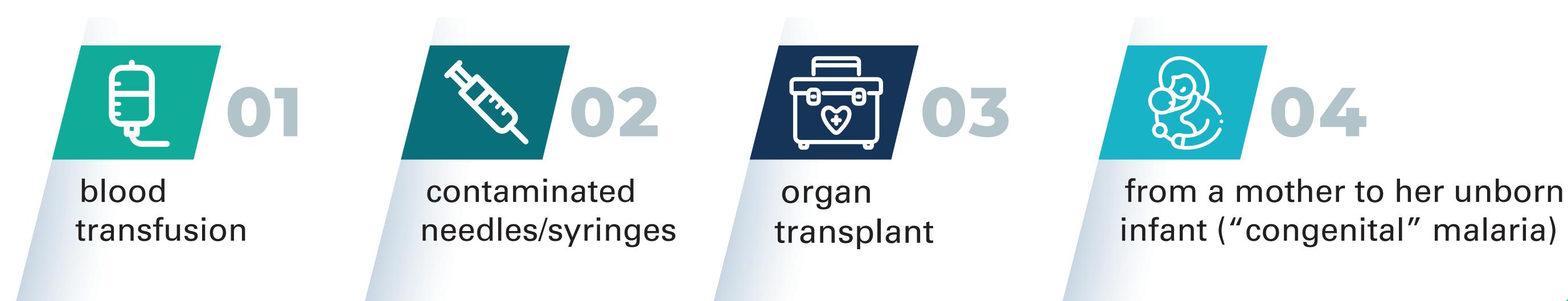
The major vector species of African malaria are members of the *Anopheles gambiae* complex and *Anopheles funestus* group.

In South Africa, the major vector species is *Anopheles arabiensis.* Potential secondary vectors are *Anopheles merus, Anopheles parensis,* and *Anopheles vaneedeni.*





IN RARE CASES, MALARIA IS TRANSMITTED THROUGH:





EOW/STEE MALARIA VECTOR CONTROLED?





One of the most effective ways to reduce malaria incidence and transmission is to reduce the population of its main vector, the Anopheles mosquito (adult and larval stages).



INDOOR RESIDUAL SPRAYING (IRS)

Many malaria vectors are considered "endophilic"; \checkmark meaning the mosquito vectors rest inside houses after taking a blood meal.

A spray technician applies residual insecticide inside a home from a rural village, Katima Mulilo, Namibia

- IRS involves coating the walls/ surfaces of a house with an insecticide.
- For several months, the insecticide will kill mosquitoes that come in contact with these surfaces.
- By killing mosquitoes after they have fed, transmission of infection to other persons is prevented.
- To be effective, IRS must be applied to a very high proportion of households in an area (usually >80%).



INSECTICIDE-TREATED BED NETS (ITN)

The use of long lasting insecticide treated nets each night is one of the most effective ways to prevent malaria, Vanuatu

- Treated bed nets form a protective barrier from mosquitoes to the people sleeping under them.
- The insecticides used for treating bed nets repel and kill mosquitoes, reducing the number that enter the house and attempt to feed on people inside.
- When high community coverage is achieved (≥50%), the number and life span of mosquitoes is reduced. This protects members of the larger community, regardless of whether they are using a bed net.



HOW STHE MALARIA VECTOR CONTROLED?



Malaria vector researchers investigate potential mosquito breeding sites for the presence of Anopheles larvae in Malahlapanga, Kruger National Park, South Africa

LARVAL SOURCE MANAGEMENT

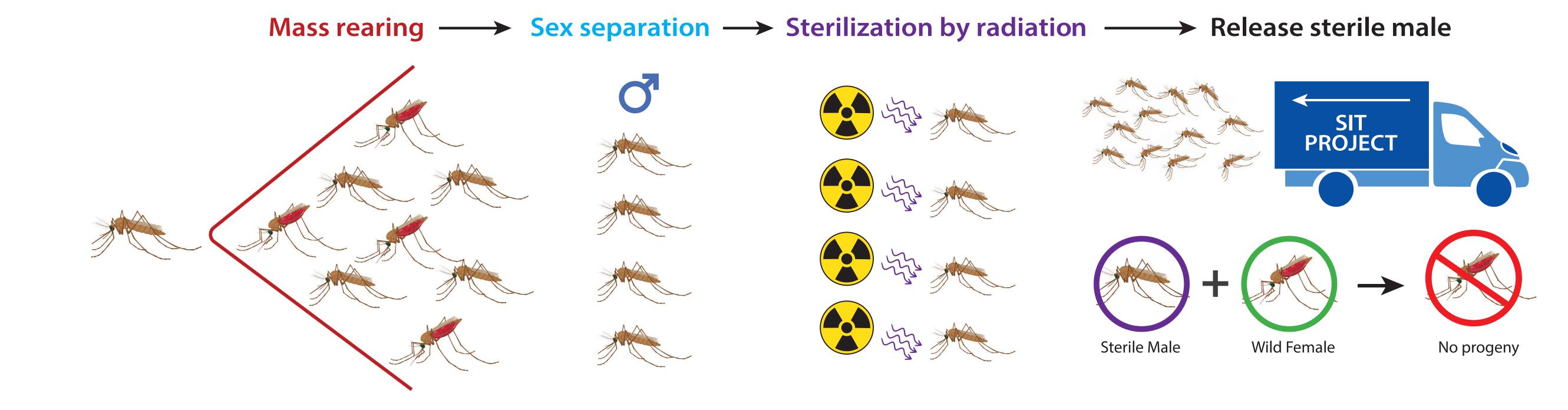
- Mosquitoes breed in water so can be controlled through careful water management.
- Larval habitats may be destroyed by filling depressions that collect water, or draining swamps or marshy areas to remove standing water.
- ✓ For some mosquito species, habitat elimination is not possible. In this case, chemical insecticides can be applied directly to the larval habitats.



NEW INNOVATIONS

Sterile Male Release

Introducing sterile male mosquitoes into an area has been successfully applied in several small-scale areas. However, the need for large numbers of mosquitoes for release makes this approach impractical for most areas.



Genetic Modification of Malaria Vectors

Genetic modification aims to develop mosquitoes that are not susceptible to the parasite. This approach is still years from application in field settings, though there have been remarkable technological advances in recent years.

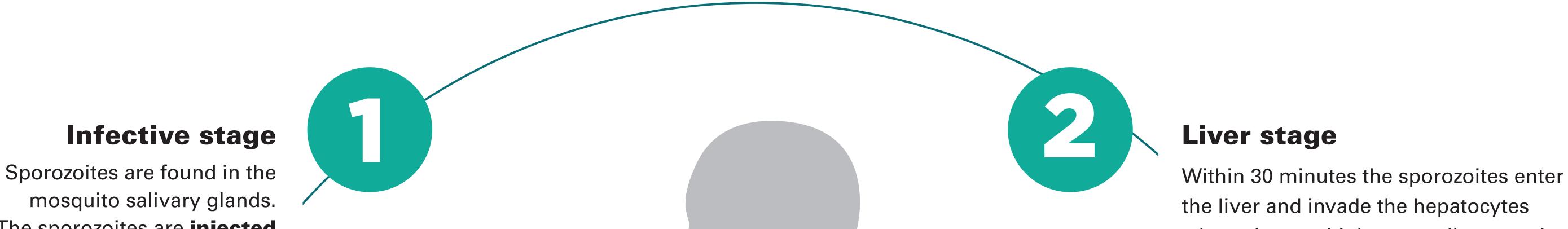




MALARA LIFE CYCLE

Malaria is a disease CAUSED BY SINGLE-CELLED PARASITES that have a life cycle REQUIRING 2 HOSTS – a mosquito vector, and a vertebrate host

5 DISTINCT LIFE CYCLE STAGES



where they multiply asexually to produce many merozoites. This asexual division continues for 7-30 days (depending on parasite species) before the merozoites are released into the blood stream. No signs or symptoms are exhibited during this stage, called the **incubation period**.

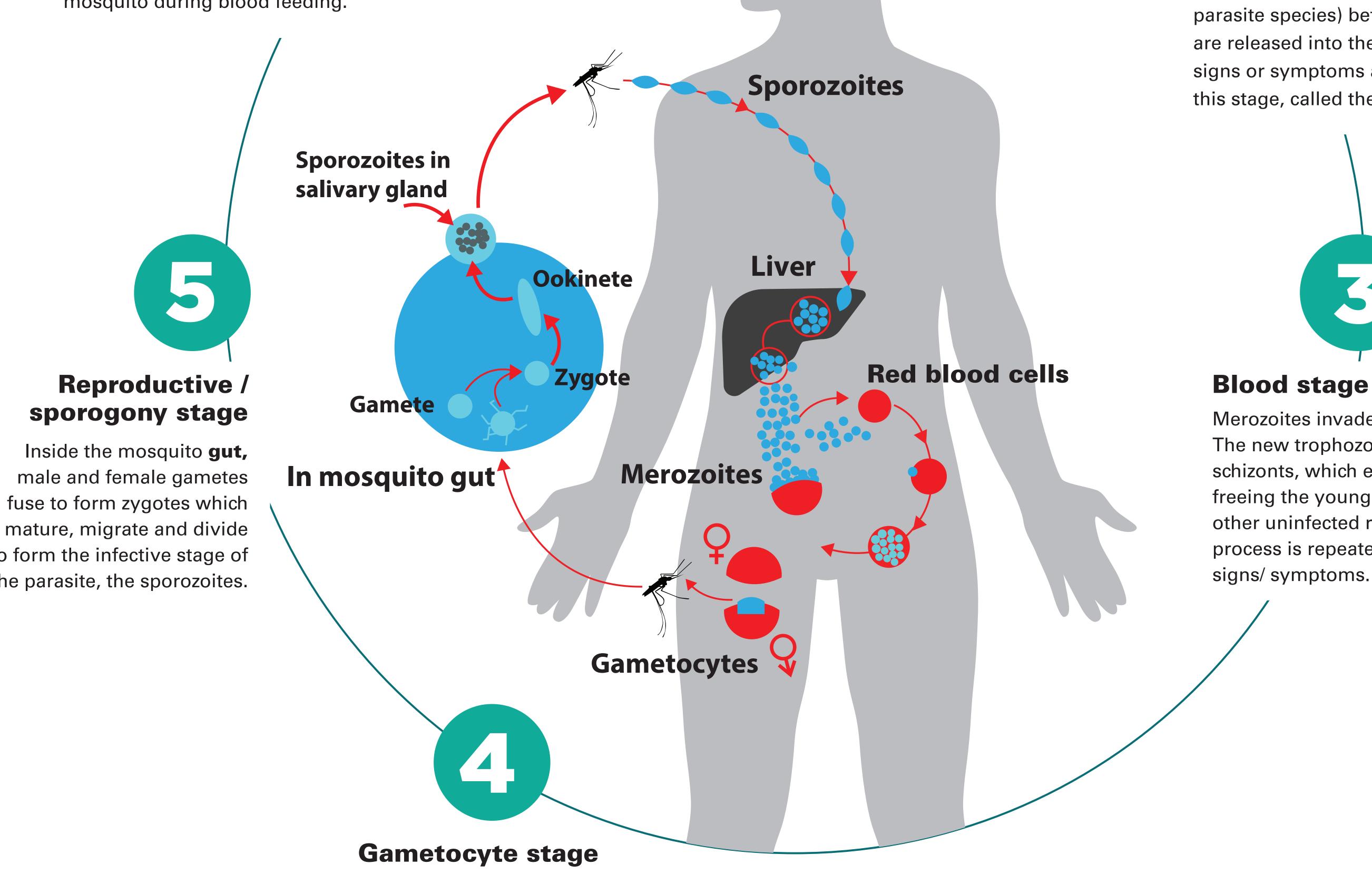


Blood stage

Merozoites invade **red blood cells**. The new trophozoites mature into schizonts, which eventually rupture freeing the young merozoites to invade other uninfected red blood cells \rightarrow this process is repeated inducing malaria

The sporozoites are **injected** into the blood stream by

an infected female Anopheles mosquito during blood feeding.



to form the infective stage of the parasite, the sporozoites.

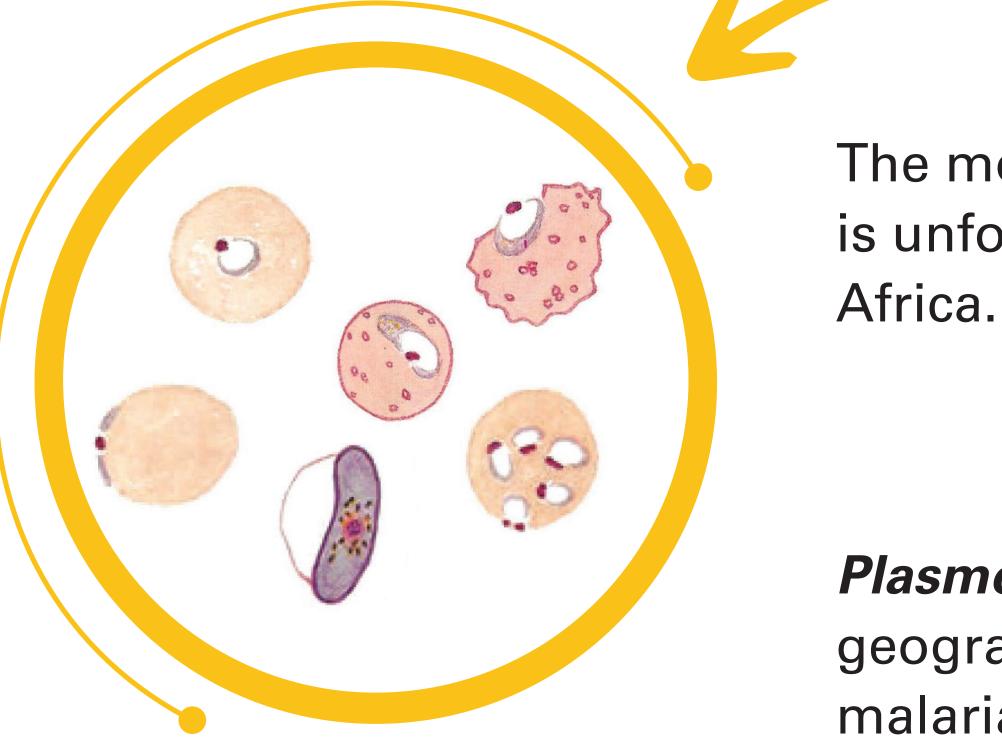
> Some parasites develop into the sexual parasite form, the gametocytes. Male and female gametocytes are taken up from the peripheral blood while a female mosquito is feeding.



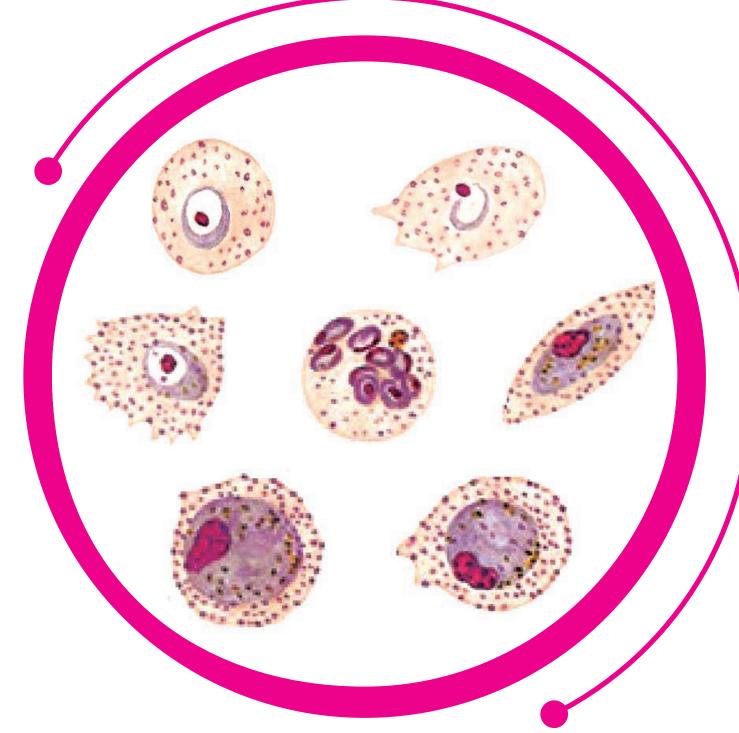
WHICH MALARIA PARASITES INFECT HUMANS?

Parasites of five different **Plasmodium** species are responsible for human malarial infections:





The most virulent, *Plasmodium falciparum*, is unfortunately also the most prevalent in

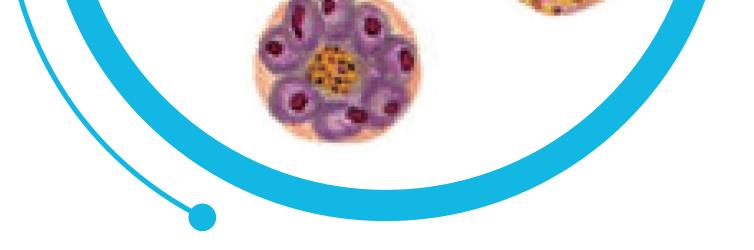


Plasmodium falciparum

Plasmodium vivax parasites have the widest geographic distribution of all the human malaria species and, together with *Plasmodium* ovale infections, are associated with relapsing (recurring) malaria.

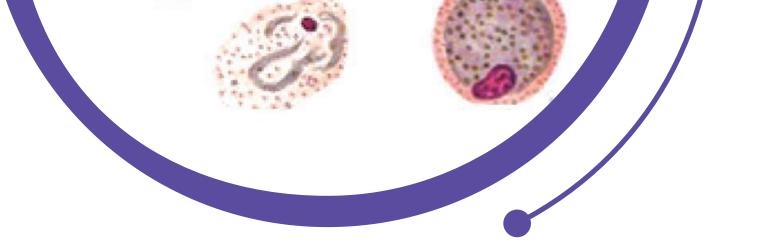
Plasmodium vivax

Under a microscope *Plasmodium malariae* and *Plasmodium knowles*i look very similar. However the disease is usually milder in Plasmodium malariae.



Plasmodium malariae

In 2010 *Plasmodium knowlesi,* a monkey malaria parasite, was recognised as the fifth human malaria parasite species following the discovery in Malaysians. It is still only reported in Southeast Asia.



Plasmodium ovale

Microscopic identification is the gold standard method for identifying the species of malaria present on stained blood films. Well-trained microscopists are able to differentiate species based on their morphological characteristics.



MALARIA PARASITE COUNTS

The most virulent malaria parasite, *Plasmodium falciparum* is unfortunately also the most prevalent in Africa.

This parasite species multiplies rapidly, invading and destroying red blood cells, causing severe anaemia. If left untreated the infection generally progresses to severe malaria, which often is fatal as it is associated with central nervous system shutdown and major organ failure.

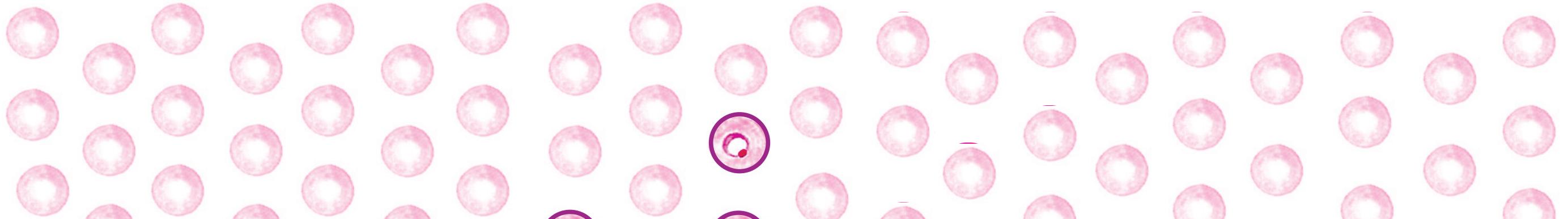


It is important to **DETERMINE THE PARASITE LOAD** with

Plasmodium falciparum infections to:

- **1.** Determine severity of infection and
- **2.** to monitor treatment efficacy.

PARASITE LOAD MAY BE EXPRESSED AS A PERCENTAGE OF INFECTED RED BLOOD CELLS OR PARASITES PER MICROLITRE OF BLOOD.



Approximately ≥5% infection is considered severe malaria, this is about 250,000 parasites per microliter of blood.



WHAT ARE THE SIGNS & SYMPTOMS OF MALARIA?

Signs and symptoms of malaria are initially **NON-SPECIFIC**.

It is **recommended that anyone with a 'flu-like illness be tested** for malaria after visiting a malaria risk area (irrespective of malaria season, transmission intensity or chemoprophylaxis use).

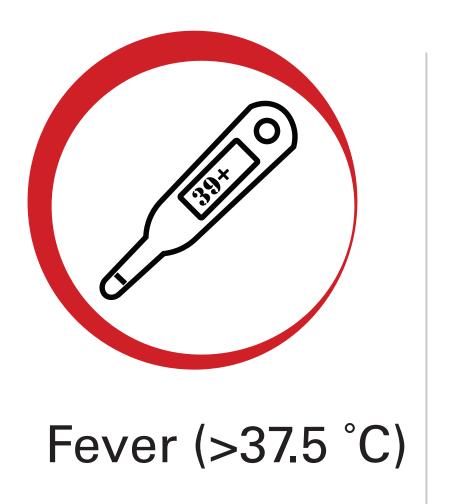
SEVERE MALARIA is a medical emergency.

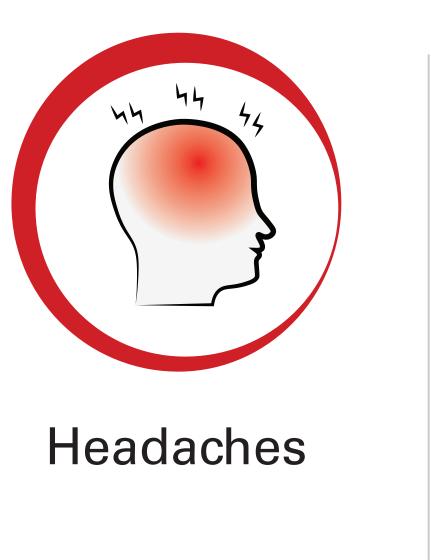
Without treatment, the clinical picture may deteriorate rapidly. Severe malaria carries a 10-40% case fatality rate in spite of treatment.



MALARIA SYMPTOMS

UNCOMPLICATED









SEVERE (COMPLICATED) MALARIA

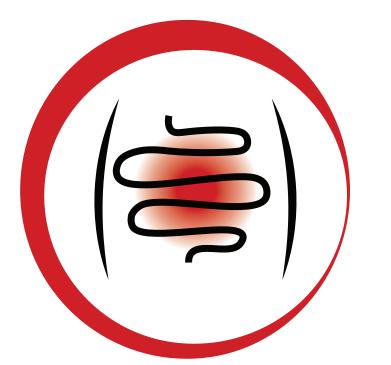
1. Impaired consciousness

2. Inability to sit or stand up straight

3. Multiple convulsions

4. Respiratory distress

5. Circulatory collapse



Diarrhoea, nausea and vomiting Loss of appetite; inability to feed in babies



Dizziness, sore throat Muscle weakness and lethargy, particularly in

young children

6. Jaundice

7. Pulmonary oedema

8. Haemoglobinuria

9. Acute respiratory distress syndrome (ARDS)



HOW DO I PREVENT AND TREAT MALARIA?

PREVENTION



Try to minimise mosquito bites, by using insect repellents, bed nets, wearing long sleeved clothes & socks after sunset.



Antimalarial medicines (chemoprophylaxis) can also be used.

Consult a health-care provider as guidelines should be followed.

TREATMENT

CONFIRM DIAGNOSIS AND ASSESS SEVERITY

Uncomplicated malaria

Mild symptoms Ambulant Normal mental function No repeated vomiting No jaundice, and no other features of severe malaria

Severe malaria

Impaired consciousness Jaundice Multiple convulsions Respiratory distress Circulatory collapse

Uncomplicated malaria caused by:

Severe malaria (usually P. falciparum)

• P. falciparum, P. malariae or P. knowlesi

artemether-lumefantrine (Coartem[®]) or, if Coartem[®] is not available, oral quinine plus either doxycycline or clindamycin

- P. ovale or P. vivax or mixed infections of P. falciparum plus P. vivax or P. ovale: artemether-lumefantrine followed by primaquine
- If unsure of species, treat as for *P. falciparum*

IV artesunate or, if not available, IV quinine. Once able to tolerate oral treatment follow with: artemether-lumefantrine (Coartem®) or, if Coartem® is not available, quinine (plus doxycycline or clindamycin)



IS THERE A VACCINE FOR MALARIA?

YES, there is one vaccine

The **RTS,S vaccine** has been approved for use; and there are many others being developed.

2019-2020

RTS,S MALARIA VACCINE EVALUATION PILOTS AND MAIN RESULTS



Significantly reduces malaria and life-threatening severe malaria (by about 30%). Since 2019, delivered



in childhood vaccination in three country-led pilots.

IN 2+ YEARS 2.4 million+ doses 830K+ CHILDREN VACCINATED

Estimated to be cost-effective in areas of moderate to high malaria transmission

30 years

of research and development

The RTS,S vaccine can be delivered through the existing platform for childhood vaccination that



On 6 October 2021, WHO recommended that the **RTS,S malaria vaccine** be used for the prevention of *P. falciparum* malaria in children living in **regions with moderate to high transmission**.

SOUTH AFRICA DOES NOT FALL IN THIS CATEGORY.



HOW IS MALARIA DIAGNOSED IN THE LAB?

electrocytosis Microcytosis Microcytosis Microcytosis

MICROSCOPY of Giemsa-stained blood films and RAPID DIAGNOSTIC TESTS (RDTs) are the routinely used methods to diagnose malaria in the laboratory.





RAPID DIAGNOSTIC TEST (RDT)

ADVANTAGES DISADVANTAGES

ADVANTAGES

DISADVANTAGES

ia Parasite Test

Quick diagnosis

- Less sensitive
- Very inexpensive •
- Parasites can be quantified
- Subjective
- Expert microscopist needed
- Very quick diagnosis
- Inexpensive
- Suitable for use in the field
- No special equipment needed
- Less sensitive
- Cannot identify
 all species
- Parasites cannot be quantified

Malaria PCR should be requested in certain instances such as when microscopy and RDT results do not correlate, microscopy and RDT are negative but malaria is still suspected or confirmation of species identification is needed.

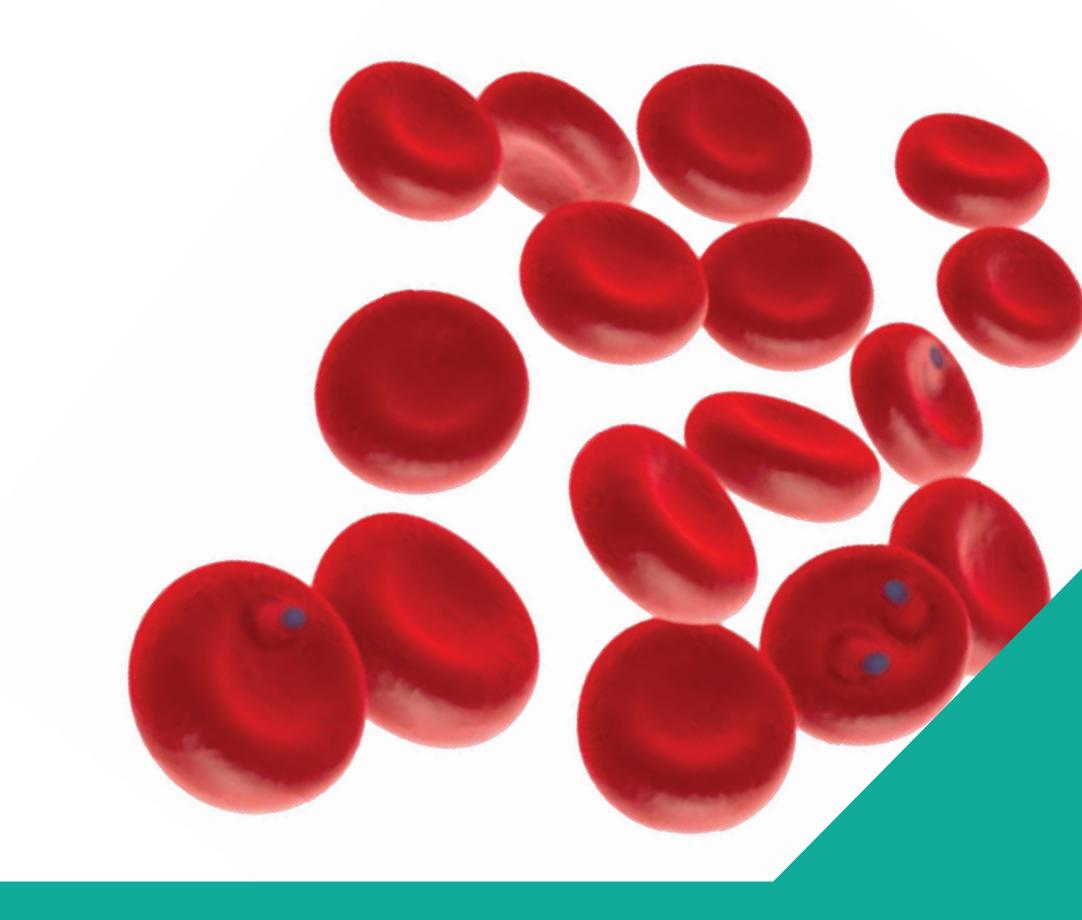
POLYMERASE CHAIN REACTION (PCR)

ADVANTAGES

- Highly sensitive
- More specific (species identification)

DISADVANTAGES

- Expensive
- Time consuming

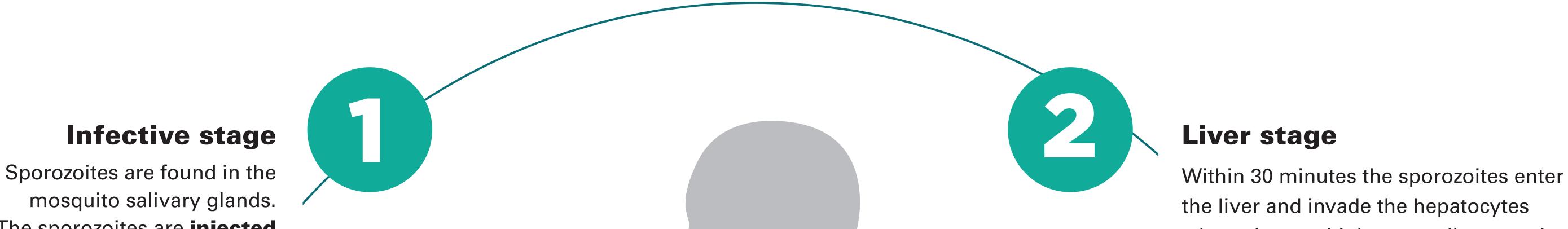




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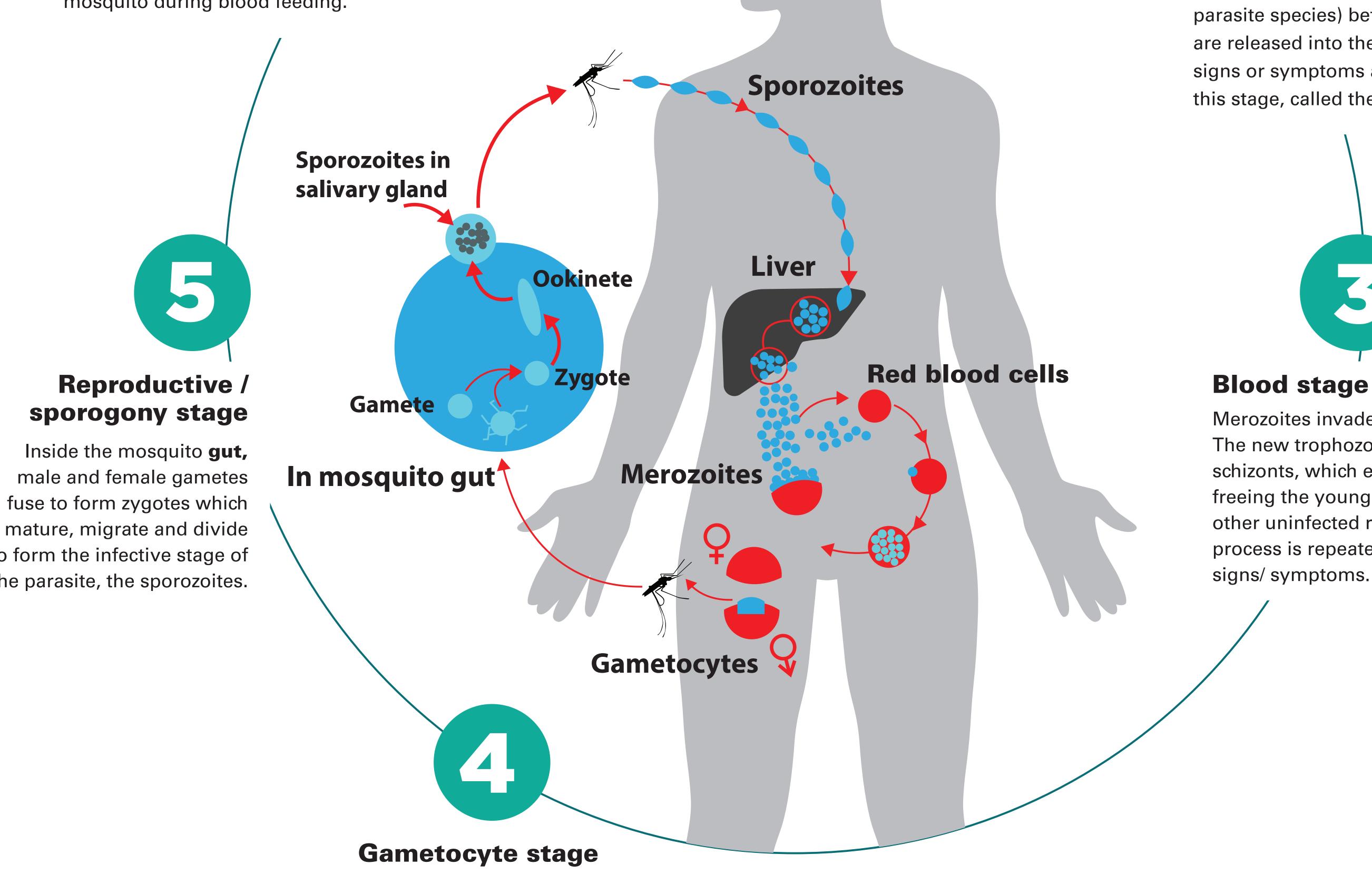


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