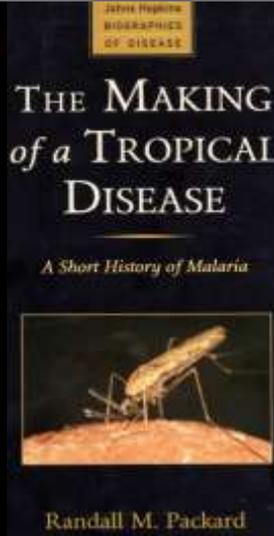


# “MALÁRIA: PERSPECTIVA HISTÓRIA”

José M. D. Poças

Médico Internista e Infeciologista



# MAKE MALARIA HISTORY

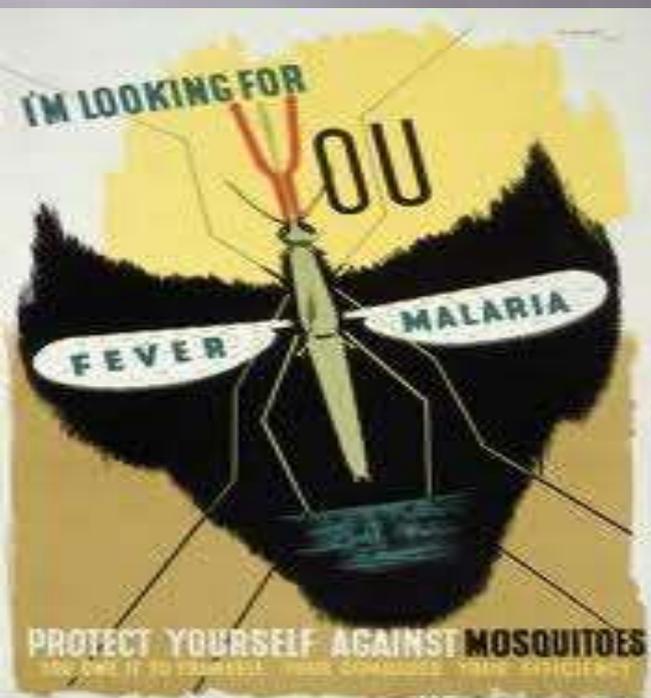
A United Methodist Campaign



DRIVE AGAINST MALARIA



## INTRODUÇÃO





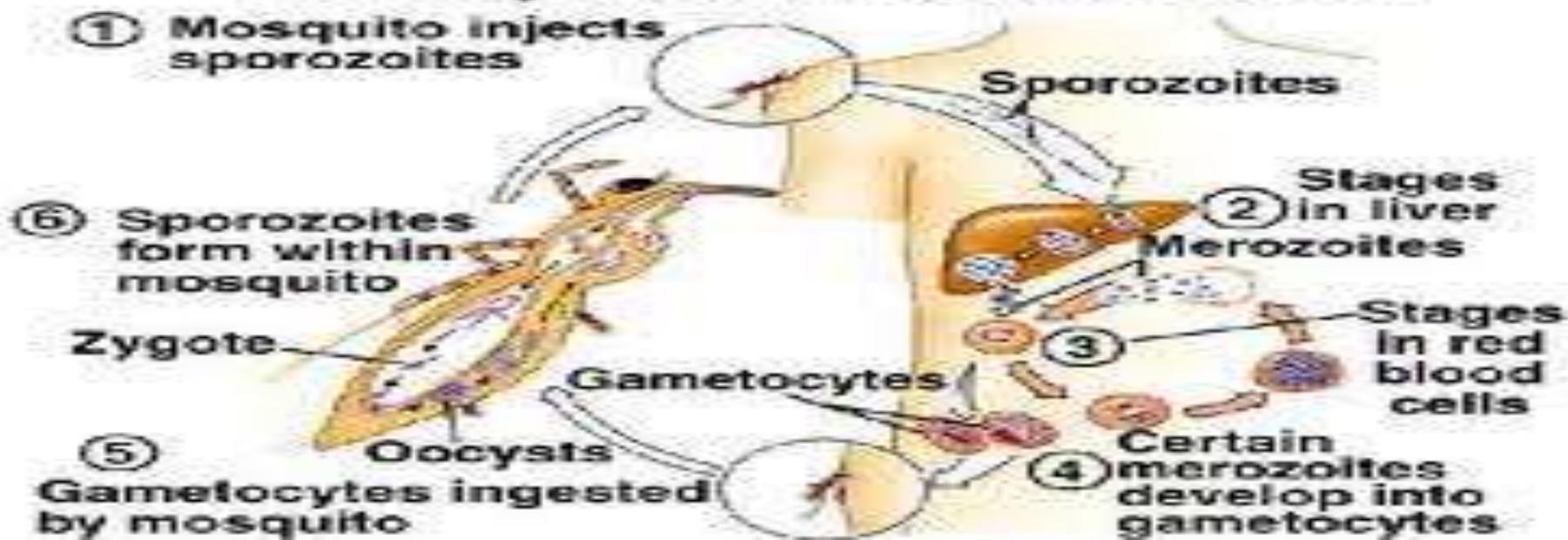
## MALARIA: Breaking the Cycle

Learn about Our  
Research Activities

[CLICK TO READ MORE >](#)

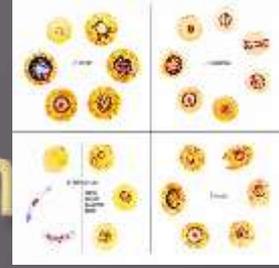
HIGHLIGHTS

## Life Cycle of *Plasmodium*



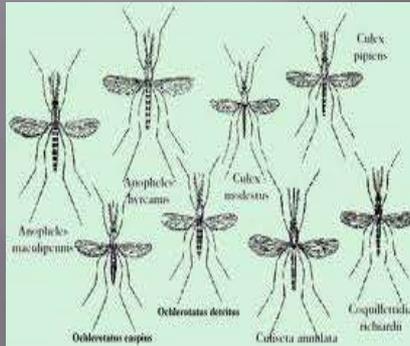


# Anopheles e Plasmodium



## Anopheles

- 460 espécies
  - 100 são transmissoras de Plasmodium aos Humanos
  - 125 demonstraram resistência aos inseticidas
  - 7 Espécies existem nos Países Mediterrâneos
    - A. Maculapennis
      - A. Atroparvus,
      - A. Claviger
      - A. Labranchiae
      - A. Messeae
      - A. Sacharovi
      - A. Hispaniola
      - A. Superpictus
- Mosquitos dos Gêneros
  - Aedes
  - Culex
  - Culiseta
  - Mansonia
  - Teobaldia
- podem também transmitir Plasmódium a outros animais, mas não aos Seres Humanos
  - Mamíferos
  - Aves
  - Répteis
- Condições Climáticas
  - -40° a 55°

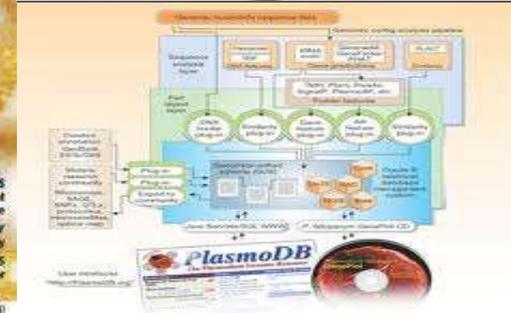


## Plasmodium

- > de 200 espécies descritas
- Cerca de 14 espécies podem infectar os Seres Humanos
  - P. falciparum
  - P. vivax
  - P. ovale curtisi e P. ovale wallikeri
  - P. malariae
  - P. knowlesi
  - P. brasilianum
  - P. cynomolgi e P. cynomolgi bastianelli
  - P. inui
  - P. rhodiani
  - P. schwetzi
  - P. semiovale

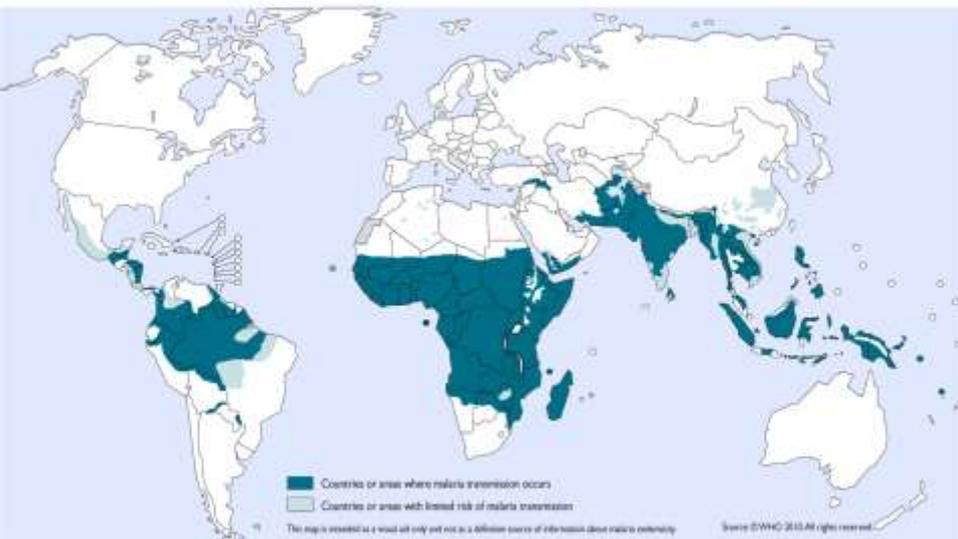


The Mosquito Genome





Malaria, countries or areas at risk of transmission, 2009



The boundaries and names shown and the designations used on this map do not imply the expression of any opinion whatsoever on the part of the World Health Organization concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. Dotted lines on maps represent approximate border lines for which there may not yet be full agreement.

Date Source: World Health Organization  
Map Production: Public Health Information and Geographic Information Systems (PHGIS)  
World Health Organization



Malaria Endemic Areas

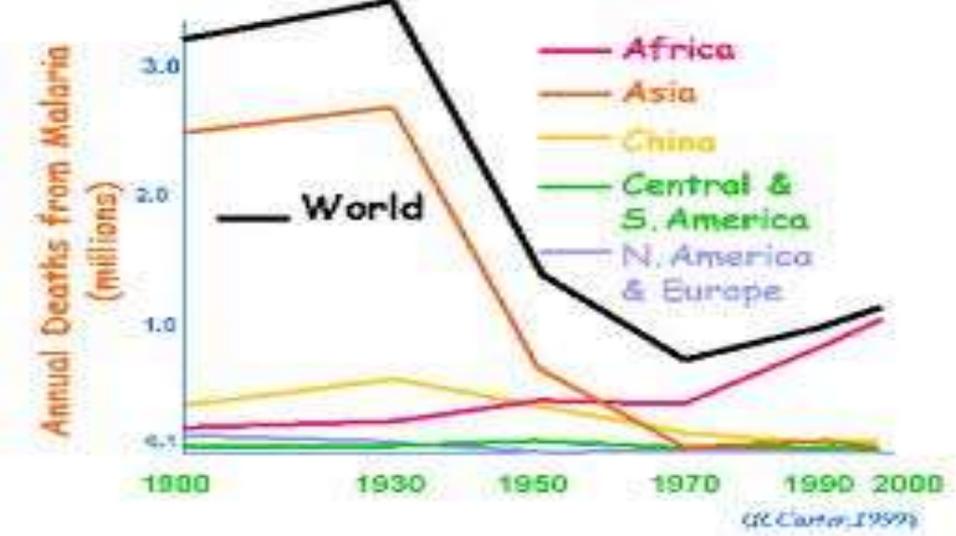
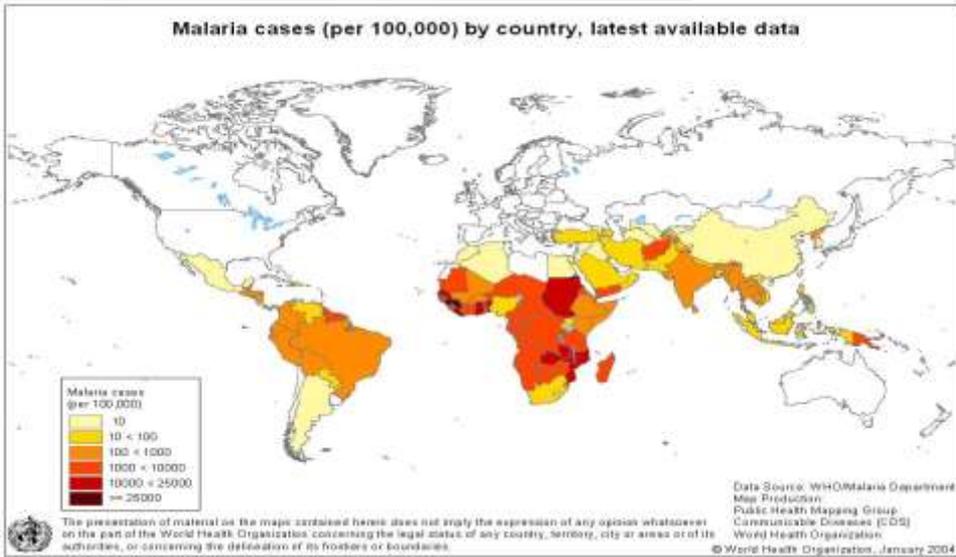


# A Child Dies EVERY 30 Seconds From MALARIA

Malaria is an economic vampire & a global threat

InfectiousBite.com

## The Global Burden of Malaria

	<i>P. falciparum</i>	<i>P. vivax</i>	<i>P. malariae</i>	<i>P. ovale</i>
Distribution	Global tropics	Global tropics and some temperate zones	Global tropics	Africa, southeast Asia, Oceania
Disease burden (number of people infected)	400 million	80-400 million	Minor	Probably minor
Number of people at risk	5 billion	3 billion	Unknown	Unknown
Type of malaria				
Asexual blood				
Asymptomatic	Many endemic settings	Many endemic settings	Almost always	Unknown
Therapies	Many	Many	Many	Many
Sexual blood				
Asymptomatic	Always	Always	Always	Always
Therapies	Primaquine, single dose	Primaquine, hyponozoitocide	Primaquine, single dose	Primaquine, hyponozoitocide
Asexual liver				
Asymptomatic	Always	Always	Always	Always
Therapies	Not indicated	Primaquine, 14 daily doses	Not indicated	Primaquine, 14 daily doses

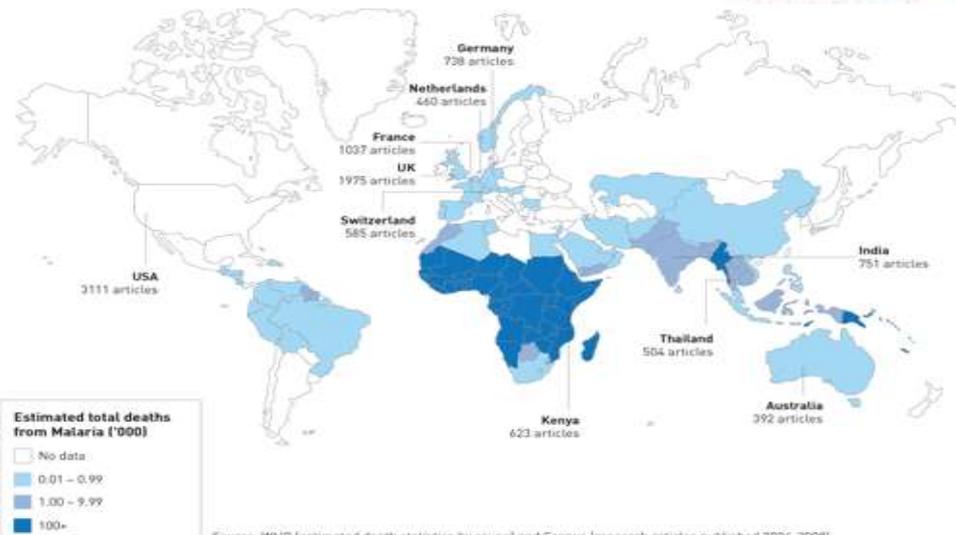


Table: Species of Plasmodium that cause malaria

# Global malaria mortality between 1980 and 2010: a systematic analysis



Christopher J L Murray, Lisa C Rosenfeld, Stephen S Lim, Kathryn G Andrews, Kyle J Foreman, Diana Haring, Nancy Fullman, Mohsen Naghavi, Rafael Lozano, Alan D Lopez

## Summary

**Background** During the past decade, renewed global and national efforts to combat malaria have led to ambitious goals. We aimed to provide an accurate assessment of the levels and time trends in malaria mortality to aid assessment of progress towards these goals and the focusing of future efforts.

**Methods** We systematically collected all available data for malaria mortality for the period 1980–2010, correcting for misclassification bias. We developed a range of predictive models, including ensemble models, to estimate malaria mortality with uncertainty by age, sex, country, and year. We used key predictors of malaria mortality such as *Plasmodium falciparum* parasite prevalence, first-line antimalarial drug resistance, and vector control. We used out-of-sample predictive validity to select the final model.

**Findings** Global malaria deaths increased from 995 000 (95% uncertainty interval 711 000–1 412 000) in 1980 to a peak of 1 817 000 (1 430 000–2 366 000) in 2004, decreasing to 1 238 000 (929 000–1 685 000) in 2010. In Africa, malaria deaths increased from 493 000 (290 000–747 000) in 1980 to 1 613 000 (1 243 000–2 145 000) in 2004, decreasing by about 30% to 1 133 000 (848 000–1 591 000) in 2010. Outside of Africa, malaria deaths have steadily decreased from 502 000 (322 000–833 000) in 1980 to 104 000 (45 000–191 000) in 2010. We estimated more deaths in individuals aged 5 years or older than has been estimated in previous studies: 435 000 (307 000–658 000) deaths in Africa and 89 000 (33 000–177 000) deaths outside of Africa in 2010.

**Interpretation** Our findings show that the malaria mortality burden is larger than previously estimated, especially in adults. There has been a rapid decrease in malaria mortality in Africa because of the scaling up of control activities supported by international donors. Donor support, however, needs to be increased if malaria elimination and eradication and broader health and development goals are to be met.

*Lancet* 2012; 379: 413–31

See Editorial page 385

Institute for Health Metrics and Evaluation, University of Washington, Seattle, WA, USA (Prof C J L Murray MD, L C Rosenfeld AB, S S Lim PhD, K G Andrews AB,

K J Foreman MPH, D Haring BSc, N Fullman MPH, M Naghavi MD, Prof R Lozano MD); and University of Queensland, School of Population Health, Herston, QLD, Australia (Prof Alan D Lopez PhD)

Correspondence to: Prof Christopher J L Murray, Institute for Health Metrics and Evaluation, 2301 Fifth Avenue, Suite 600, Seattle, WA 98122, USA (cjm@u.washington.edu)

	1980–89	1990–99	2000–10	Total
Vital registration (national)	262	340	313	915
Vital registration (subnational)	15	7	2	24
Verbal autopsy (national)	0	2	12	14
Verbal autopsy (subnational)	41	44	64	149
Other	0	27	21	48

Table 1: Site-years of data, by decade

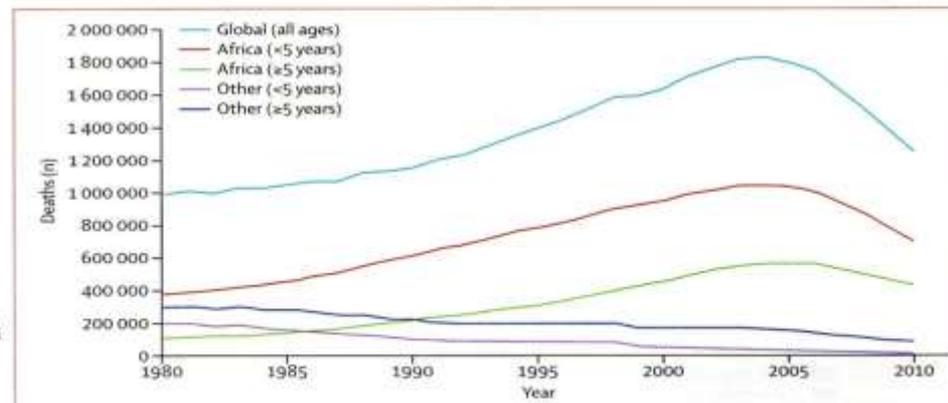


Figure 2: Trends in global malaria deaths by age and geographical region, 1980 to 2010

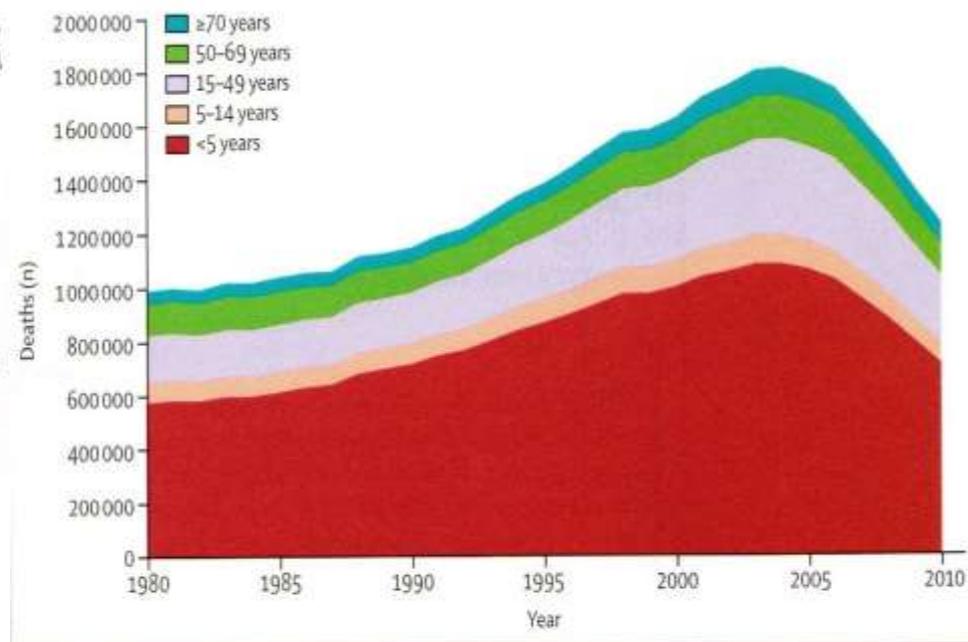
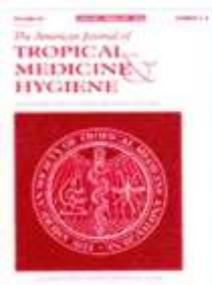


Figure 6: Malaria deaths by age, 1980 to 2010



## The Intolerable Burden of Malaria: A New Look at the Numbers

Supplement to Volume 64(1) of the *American Journal of Tropical Medicine and Hygiene*

Edited by Joel G Breman, Andréa Egan, and Gerald T Keusch.

Fogarty International Center, National Institutes of Health, Bethesda, Maryland; Multilateral Initiative on Malaria, Fogarty International Center, National Institutes of Health, Bethesda, Maryland

Northbrook (IL): American Society of Tropical Medicine and Hygiene, January 2001.

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## The Intolerable Burden of Malaria: A New Look at the Numbers

Joel G. Breman, Andréa Egan, and Gerald T. Keusch.

Fogarty International Center, National Institutes of Health, Bethesda, Maryland; Multilateral Initiative on Malaria, Fogarty International Center, National Institutes of Health, Bethesda, Maryland

*"When you cannot measure it, when you cannot express it in numbers, you have scarcely . . . advanced to the stage of Science, whatever the matter may be."*

William Thomson, Lord Kelvin, 1824–1907

## The Economic Burden of Malaria

John Luke Gallup\* and Jeffrey D. Sachs.

Center for International Development, Harvard University, Cambridge, Massachusetts

### Abstract

Malaria and poverty are intimately connected. Controlling for factors such as tropical location, colonial history, and geographical isolation, countries with intensive malaria had income levels in 1995 of only 33% that of countries without malaria, whether or not the countries were in Africa. The high levels of malaria in poor countries are not mainly a consequence of poverty. Malaria is geographically specific. The ecological conditions that support the more efficient malaria mosquito vectors primarily determine the distribution and intensity of the disease. Intensive efforts to eliminate malaria in the most severely affected tropical countries have been largely ineffective. Countries that have eliminated malaria in the past half century have all been either subtropical or islands. These countries' economic growth in the 5 years after eliminating malaria has usually been substantially higher than growth in the neighboring countries. Cross-country regressions for the 1965–1990 period confirm the relationship between malaria and economic growth. Taking into account initial poverty, economic policy, tropical location, and life expectancy, among other factors, countries with intensive malaria grew 1.3% less per person per year, and a 10% reduction in malaria was associated with 0.3% higher growth. Controlling for many other tropical diseases does not change the correlation of malaria with economic growth, and these diseases are not themselves significantly negatively correlated with economic growth. A second independent measure of malaria has a slightly higher correlation with economic growth in the 1980–1996 period. We speculate about the mechanisms that could cause malaria to have such a large impact on the economy, such as foreign investment and economic networks within the country.

# THE GLOBAL MALARIA BUSINESS PLAN (Executive Summary)

OF THE



# RBM

ROLL BACK MALARIA PARTNERSHIP

MAY 5, 2008

**PRELIMINARY DRAFT FOR DISCUSSION**





# Malaria Site

all about malaria

## Time Line For Origin of Malaria

Half a billion years ago	Existence of pre-parasitic ancestor
150 million to 200 million years ago	Early Dipterans, ancestors of mosquitoes, appear
130 million years ago	Two-host life cycle in Dipterans and vertebrates evolves
130 million years ago	Divergence of the bird and mammalian malaria parasites
100 million years ago	Lineage of <i>P. malariae</i> , <i>P. ovale</i> , and <i>P. vivax</i> diverges
~5 million years ago	<i>P. falciparum</i> evolves
2-3 million years ago	Divergence of <i>P. vivax</i> from <i>P. cynomolgi</i>
4000-10000 years ago	Lethal strain of <i>P. falciparum</i> appears
4000-5000 years ago	<i>Anophelines</i> in Africa develop highly anthropophilic habits

## Time Line For Spread of Malaria

>10000 years ago	Malaria in Africa
10000-5000 years ago	Mesopotamia, the Indian peninsula and South-East Asia
5000 years ago	Malaria in China
3000 years ago	<i>P. falciparum</i> reaches India
2,500 - 2,000 years ago	Malaria reaches the Mediterranean shores
1000-500 years ago	Malaria reaches northern Europe
End of 15th century AD	Malaria reaches New World
Mid 18th century AD	Malaria spreads across North America
19th Century AD	Malaria almost all over the globe
Early 20th Century AD	Millions die of malaria almost all over the world
Early 1950s	Malaria almost disappears from North America and from almost all of Europe; deaths mainly in Africa

# Evolução das Hemoglobinopatias (10.000 - 20.000 anos)

- Mutações “Protectoras”
  - Talassémias
  - G6PH (deficiência em)
  - Drepanocitose
  - Hemoglobina C
  - Hemoglobina E
  - Ovalovitose
- Outros Factores
  - Grupos Sanguíneos (DuffyNegativo)

CLINICAL MICROBIOLOGY REVIEWS, Oct. 2002, p. 564-594  
0893-8512/02/\$04.00-0 DOI: 10.1128/CMR.15.4.564-594.2002  
Copyright © 2002, American Society for Microbiology. All Rights Reserved.

Vol. 15, No. 4

## Evolutionary and Historical Aspects of the Burden of Malaria

Richard Carter<sup>1\*</sup> and Kamini N. Mendis<sup>2</sup>

*University of Edinburgh, Division of Biological Sciences, ICAPB, Ashworth Laboratories, Edinburgh EH9 3JT, United Kingdom,<sup>1</sup> and Roll Back Malaria Project, World Health Organization, Geneva 27, Switzerland<sup>2</sup>*



## Malaria and the Red Cell

*David J. Weatherall, Louis H. Miller, Dror I. Baruch, Kevin Marsh, Ogobara K. Doumbo, Climent Casals-Pascual, and David J. Roberts*

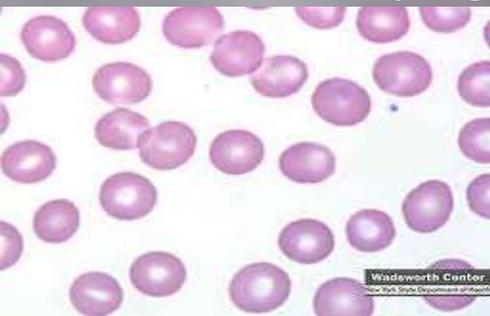
Because of the breakdown of malaria control programs, the constant emergence of drug resistant parasites, and, possibly, climatic changes malaria poses a major problem for the developing countries. In addition, because of the speed of international travel it is being seen with increasing frequency as an imported disease in non-tropical countries. This update explores recent information about the pathophysiology of the disease, its protean hematological manifestations, and how carrier frequencies for the common hemoglobin disorders have been maintained by relative resistance to the malarial parasite.

In Section I, Dr. Louis Miller and colleagues consider recent information about the pathophysiology

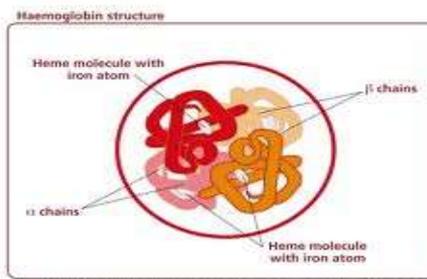
of malarial infection, including new information about interactions between the malarial parasite and vascular endothelium.

In Section II, Dr. David Roberts discusses what is known about the complex interactions between red cell production and destruction that characterize the anemia of malaria, one of the commonest causes of anemia in tropical countries.

In Section III, Dr. David Weatherall reviews recent studies on how the high gene frequencies of the thalassemias and hemoglobin variants have been maintained by heterozygote advantage against malaria and how malaria has shaped the genetic structure of human populations.



Wadsworth Center  
New York State Department of Health



# PALUDISMO / MALÁRIA

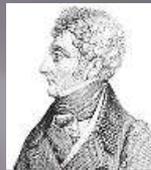
## ▣ Sinónimos

- Seção (Séc XIII)
  - Sezonismo (Ricardo Jorge: 1901)
- Malária (G. Risorì)
  - “Mau ar” (Itália Séc. XVIII)
  - “Mãe das Febres”
- Paludismo (E. Marchiafava e A. Celli)
  - “Paul, Palude” (Itália Séc. XIX)
  - Febre Palustre

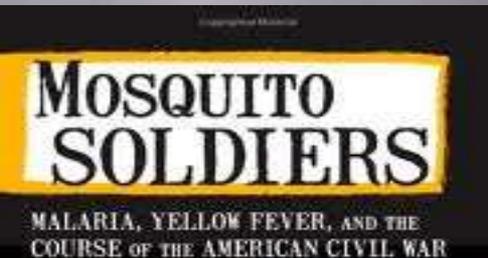


## ▣ Guerras

- Guerra Civil dos EUA
  - 1.316.000 casos
  - 10.000 mortos
- I Guerra Mundial
  - Tropas Inglesas
    - 124.000 casos
    - 23.762 mortos
- II Guerra Mundial
  - Tropas Americanas
    - 60.000 mortos
- Guerra da Coreia
  - Tropas Americanas
    - 3.000 casos
    - 629 mortos
- Guerra do Vietnam
  - Tropas Americanas
    - 40.000 casos
    - 78 mortes



Personnel of the 17th Malaria Survey Unit, Oua Bay, New Guinea, July 1963



ANDREW McILWAINE BELL





# Os que Adoeceram



## ▣ Algumas Personagens Importantes

1. REIS E IMPERADORES (BELISARIUS, KANGXI, LUISA MARIA, JAIME I, CARLOS II, CARLOS V, HANNIBAL, ISABEL, FILIPE II, FILIPE IV, FILIPE V, FERNANDO VI)
2. PRESIDENTES DOS EUA (GEORGE WASHINGTON, JAMES MONROE, ANDREW JACKSON, ULYSSES GRANT, JAMES GARFIELD, THEODORE ROOSEVELT, JOHN KENNEDY)
3. OUTROS (HOMERO, HIPOCRATES, CRISTOVÃO COLOMBO, ALBERT DURER, WILLIAM SHAKESPEARE, SIR ARTHUR CONAN DOYLE, ALFRED RUSSELL WALLACE, MERIWETHER LEWIS, MORTON STANLEY, JEFFERSON DAVIS, LUCRETIA GARFIELD, GEORGE MCCLELLAN, HO CHI MINH, JESSE JAMES, JOHN PERSHING, MAHATMA GANDHI, ERNEST HEMINGWAY, LORD NELSON, LEON TROTSKY, EUGENE O'NEILL, SIR HARRY SECOMBE, ROSS KEMP, SANTA TERESA DE JESUS, HERNAN CORTÉS, DON ADAMS, ERROL FLYNN, PETA WILSON, CAROL LANDIS, RAYMOND BURR, AIDIE MURPHY, MICHAEL CAINE, CHRISTOPHER LEE, MICHAEL DIDIKOFF, JEREMY PIVEN, AL JOHNSON, JANE GOODALL, DAVY CROCKETT, STEVE REEVES, CHRIS MATTHEWS, ANDERSON COOPER, WILSON KIPETER, YAKUBU AIYEBENI, DIKEMBE MUTAMBOI, EZEKIEL KEMBOI, MADRE TERESA DE CALCUTÁ, LEANDER PAES, ETC)



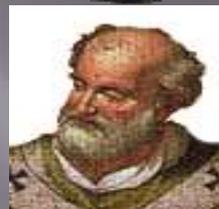


# Os que Morreram



## ALGUMAS PERSONAGENS IMPORTANTES

- Santos (Santo Agostinho)
- Reis e Imperadores (, Cesar Augustus , Alexandre da Macedónia; Alarico, Otão I e Otão II; Frederico I e Frederico IV; Andronicus Palaeopologus Henrique do Luxemburgo, Genghis Khan, Sultão Muhammed Tughluk, Eduardo VI, Minas, Mongkut)
- Guerreiros e Militares(Atila, Belisarius, Francis Ona, Comodoro Olivier Hazard Perry)
- Nobres (Richard of Cornwall, Cesare Borgia)
- Papas (Alexandre VI; Gregorio V, Damasus II, Leão X, Sistus V, Urbano VII, Gregorio XV)
- Escritores e Artistas (Petrarca, Dante, Caravaggio, Guido Cavalcanti, Lord Byron, Amrish Puri)
- Exploradores (Alvaro Mendana, David Livingstone)
- Inventores (Josep Ressel)
- Atletas (Rebka Chenashu, Didier Zohora)



# History of Malaria Parasite and its Global Spread

## Time Line for Origin of Malaria

Half a billion years ago  
Existence of pre-parasitic ancestor  
150 million to 200 million years ago  
Early Dipterans, ancestors of mosquitoes,  
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130 million years ago  
Two-host life cycle in Dipterans and  
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130 million years ago  
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Malaria almost disappears from North  
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deaths mainly in Africa

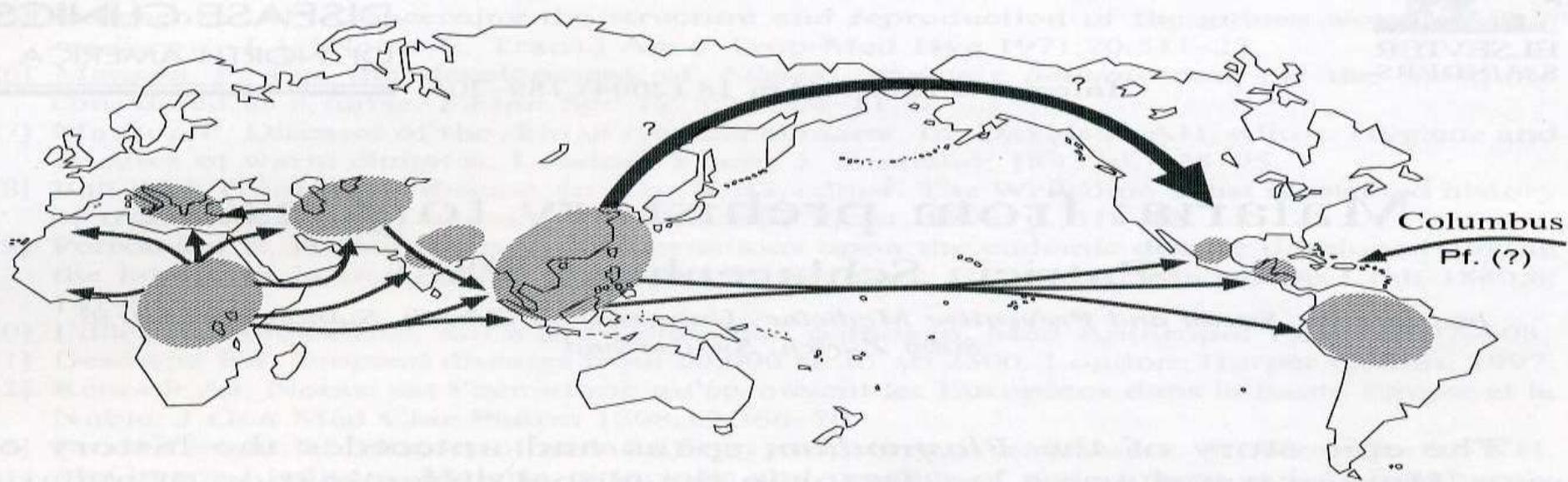


Fig. 1. Probable routes of the spread of malaria in prehistoric and early historic times. (Adapted from Bruce-Chwatt LJ. Paleogenesis and paleo-epidemiology of primate malaria. Bull World Health Organ 1965;2:363-87; with permission.)



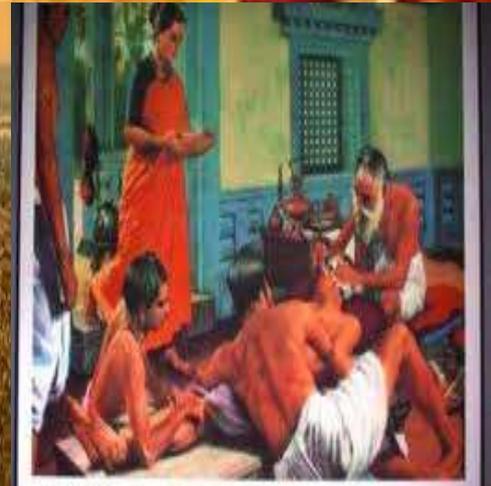
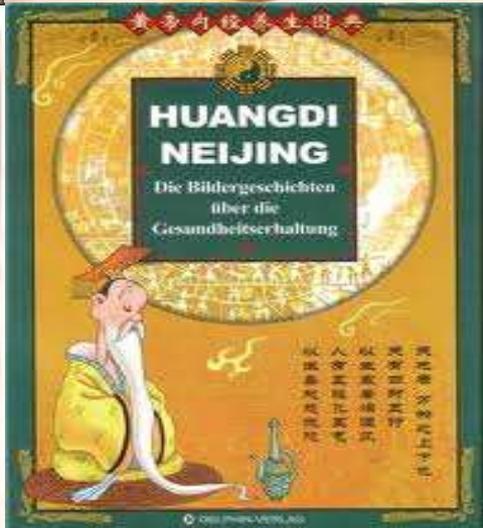


# History of Malaria: Scientific Discoveries

Time Line For Scientific Discoveries	
Ancient Times	160-200 AD
Early man attributed the fevers to evil spirits, angered deities, demons, or the black magic of sorcerers	Galen suggested that malaria was due to a disorder in the four humors of the body and suggested bleeding and/or purging as treatment; this view stayed for 1500 years
Several thousand years ago	1696 AD
Babylonian cuneiform script attributes malaria to a god, pictured as a mosquito-like insect	Morton presented the first detailed description of the clinical picture of malaria and its treatment with cinchona.
800 BC	1712 AD
Indian sage Dhanvantari wrote that bites of mosquitoes could causes diseases, fever, shivering etc.	Fransesco Torti accurately described the intricate course of the disease that was curable by the cinchona
400 BC	1716-1717 AD
Hippocrates described the various malaria fevers of man; distinguished the intermittent malarial fever from the other continuous fevers; noted the daily, every-other-day, and every-third-day temperature rise; mentioned about splenic change in malaria; attributed malaria to ingestion of stagnant water; also related the fever to the time of the year and to where the patients lived.	Lancisi first described a characteristic black pigmentation of the brain and spleen in the victims of malaria. He linked malaria with poisonous vapours of swamps or stagnant water on the ground. In 1717, in his monograph titled Noxious Emanations of Swamps and Their Cure, he echoed the view that malaria was due to minute "bugs" or "worms" which entered the blood and revived the old idea that mosquitoes might play a role.
300 BC	1796 AD
Charaka Samhita in India classifies the fevers into five different categories, namely continuous, remittent, quotidian, tertian and quartan fevers.	John Crawford, American physician contradicted the bad-air theory and suggested that the eggs, laid during a mosquito bite, hatched in the wound and migrated through the host's body, producing the manifestations of malaria
100 BC	1816 AD
Susruta Samhita in India associates fevers with the bites of the insects	Giovanni Rasori doubted the "bad air" theory and suggested that a microorganism is responsible for the disease
First Century BC	1847 AD
Roman agriculturist Collumella suggested that the diseases are caused by animals that breed in the marshes	German physician, Heinrich Meckel, identified round, ovoid, or spindle-shaped structures containing black pigment granules in protoplasmic masses in the blood and in the spleen; he probably saw the malaria parasites for the first time, but could not recognize the true importance of his finding
First Century AD	
Roman scholar Marcus Terentius Varro suggested that the grave maladies are caused by inhalation of certain animalcula that breed in the swamps	
30 AD	
Celsus described two types of tertian fevers	



Restoration of the Dalai Lama. To quantify and qualify persons' risk and

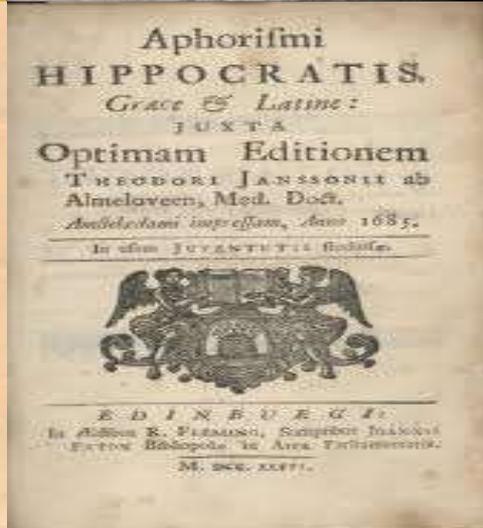
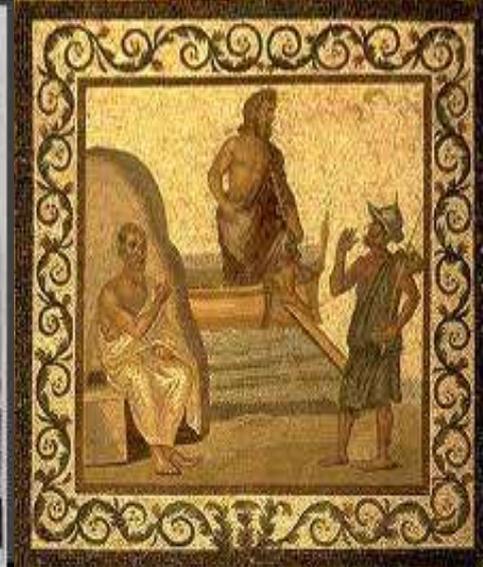
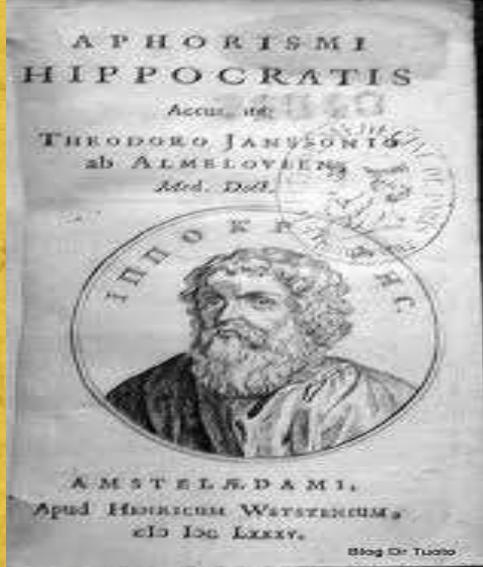
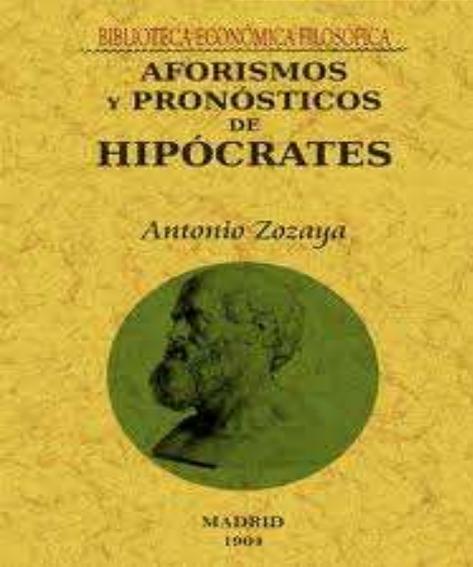


SUSRUTA : SURGEON OF OLD INDIA



"Should there be rivers in the land, which drain off from the ground the stagnant water and the rain water, the people will be healthy and bright. But if there be no rivers, and the water that the people drink to be marshy, stagnant, and fenny, the physique of the people must show protruding bellies and enlarged spleens."

Hippocrates



# History of Malaria: Scientific Discoveries

1848-1850 AD	1893
Schutz, Virchow and Hischl confirmed the presence of pigment with intermittent fevers.	Golgi suggested that malaria parasites may have an undiscovered tissue phase in endothelial cells not affected by antimalarial drugs and could be the source of relapses.
1850 AD	1897
American Josiah Clark Nott dismissed the miasma theory and suggested that microscopic "insects" transmitted by mosquitoes caused both malaria and yellow fever	William G. McCallum and Ople demonstrated the sexual process of the malaria parasite. On August 20, 1897, a tired, discouraged Ronald Ross
1854 AD	August 20, 1897
Beauperthy, American naturalist, wrote that malaria and yellow fever "are produced by venomous fluid injected under the skin by mosquitoes	Ronald Ross demonstrated oocysts in the gut of anopheline mosquito at Secunderabad, India, proving that mosquito was the vector for malaria
1878-79 AD	July, 1898
Edwin Klebs and Corrado Tommasi-Crudeli announced the "discovery" of <i>Bacillus malariae</i> , a bacteria that supposedly caused malaria	Ronald Ross demonstrated the sporozoites in the salivary glands of the mosquito and then transmitted malaria to birds through infected mosquitoes.
November 6, 1880	September, 1898
Charles Louis Alphonse Laveran, a French physician working in Algeria, found a moving object while examining a fresh blood film from a patient of malaria. He called this parasite <i>Oscillaria malariae</i> .	Giovanni Battista Grassi transmitted vivax malaria to a human volunteer
1881	1900
George Sternberg, American bacteriologist proved that the <i>Bacillus malariae</i> was not responsible for malaria	Patrick Manson transmitted malaria to volunteers in London from infected mosquitoes brought from Italy.
1882	1911
Albert Freeman Africanus King, a gynecologist at George Washington University, suggested that the mosquito is the real source of malaria	Brown suggested the hematin origin of the 'black pigment' and suggested the action of a proteolytic enzyme on hemoglobin to be the most probable mode of elaboration of the malaria pigment
1884	1948
Russian physiologist, Basil Danielewsky identified malaria parasites of in the blood of wild birds	Shortt & Garnham demonstrated the tissue form of <i>P. vivax</i> malaria
1884	1975
Marchiafava and Celli demonstrated active amoeboid ring in unstained blood and named it <i>Plasmodium</i> ,	William Trager cultured <i>P. falciparum</i> inside red blood cells
1886	1977
Louis Pasteur, William Osler and Camillo Golgi confirm Laveran's finding	Lysenko et al. suggested latent forms of <i>P. vivax</i> malaria that cause relapses
1886	1987
Pel suggested the existence of a tissue stage of the parasite	Dr. Manuel Elkin Patarroyo, a Colombian biochemist, developed the first synthetic Spf66 vaccine against <i>P. falciparum</i>
1886	2002
Golgi observe that the tertian and quartan forms produced differing numbers of segmentations on maturity; also demonstrated that the fever coincided with the rupture and release of merozoites into the blood stream	The genome of <i>Anopheles gambiae</i> and <i>Plasmodium falciparum</i> sequenced
1891	2008
Romanowsky described better staining methods for identifying malarial parasites	The genome of <i>P. vivax</i> and <i>P. knowlesi</i> sequenced

# Alphonse Laveran's discovery 100 years ago and today's global fight against malaria<sup>1</sup>

L J Bruce-Chwatt CMG OBE MD FRCP

Wellcome Museum of Medical Science, London NW1

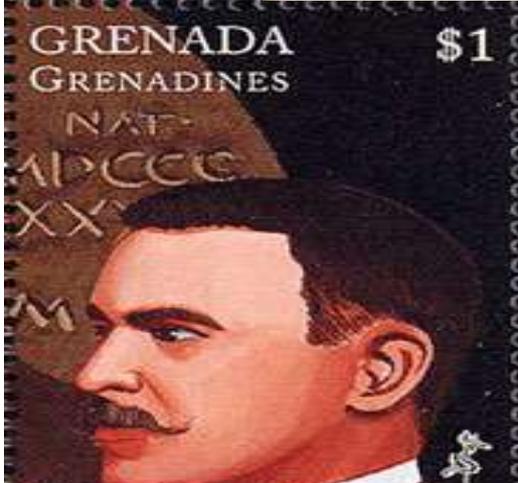
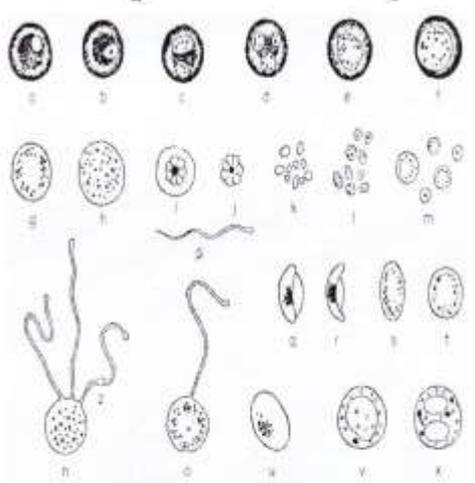
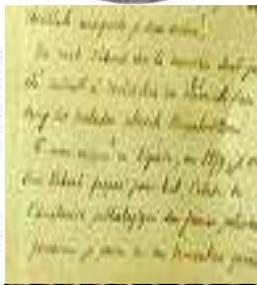
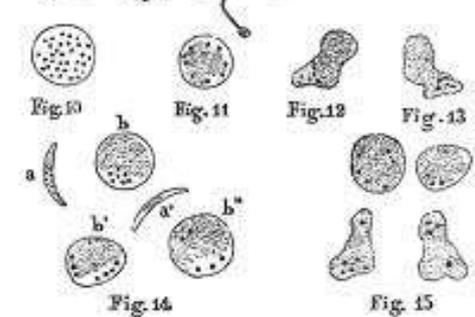
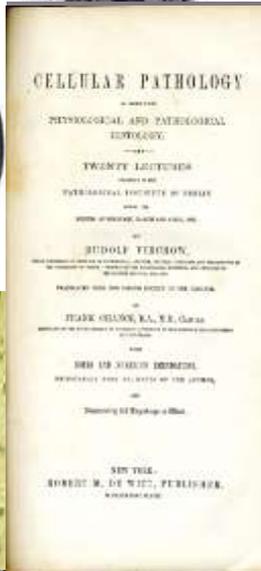
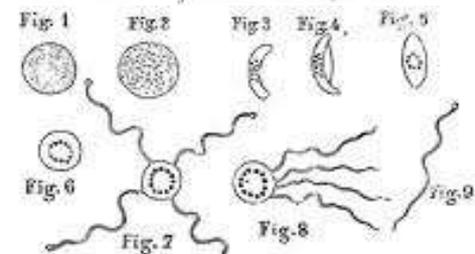
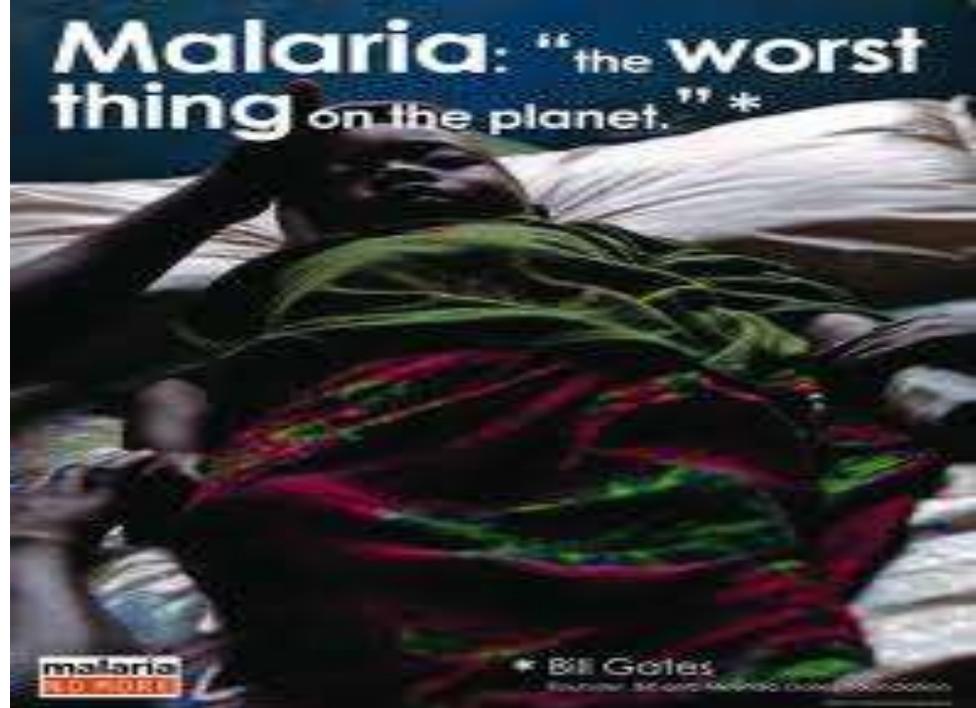
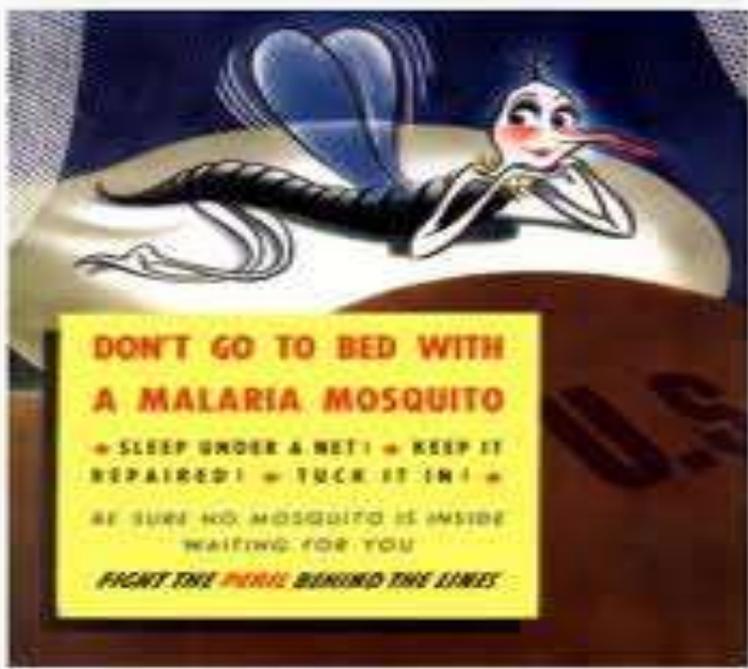
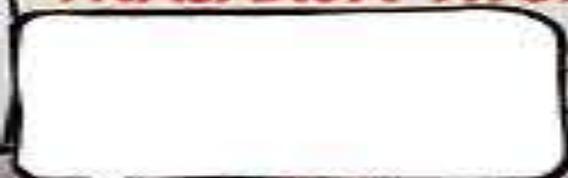


Figure 1. Malaria parasites (*Haemamoeba malariae*) observed and drawn by Alphonse Laveran; published in *Comptes Rendus de l'Academie des Sciences*, 24 October 1881. (From Sergent & Sergent 1929)

Sir Ronald Ross (1858-1930) *Physiology of Medicine*, 1902



## MALARJA THEME PARTY MUMBAI



# History of Malaria Treatment

## Time Line For Malaria Treatment

340 AD

Anti-fever properties of *qinghao* first described by Ge Hong in China

Between 1620 and 1630 AD

Spanish Jesuit missionaries in Peru learn the healing power of a tree bark

1632

Jesuit Barnabé de Cobo takes Cinchona bark to Europe

1633

Properties of the bark in the treatment of malaria first written by Father Antonio de la Calancha

1640

Juan de Lugo first employed the tincture of the cinchona bark for treating malaria in Countess of Chinchon

1658

The first prescription of cinchona in England by Robert Brady

1670s

Robert Talbor develops an infusion of cinchona powder in white wine and uses it as a 'secret remedy'

1712

Fransesco Torti writes a book on the therapeutic properties of the bark

1742

Linnaeus, a Swedish botanist, classifies the Peruvian bark and names the tree *cinchona*

1820

French chemists Joseph Pelletier and Jean Biename Caventou isolate quinine

1844/1910

Sporadic resistance to quinine reported

1934

Resochin (chloroquine) synthesised at Bayer, Germany by Hans Andersag

1944

Proguanil or Paludrine (chlorguanide hydrochloride) developed by Curd, Davey and Rose

1945, 1950

Camoquin (amodiaquin) and Primaquine (Elderfield) developed

1952

Pyrimethamine developed

1971

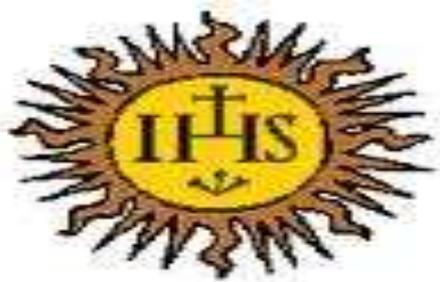
The active ingredient of *qinghao* isolated by Chinese scientists

1974-75

Mefloquine jointly developed by the U.S. Army Medical Research and Development Command, World Health Organization and Hoffman-La Roche

1992 and 1998

Atovaquone becomes available in 1992; a combination of proguanil and atovaquone, called Malarone, becomes available in Australia in 1998



The English Connection

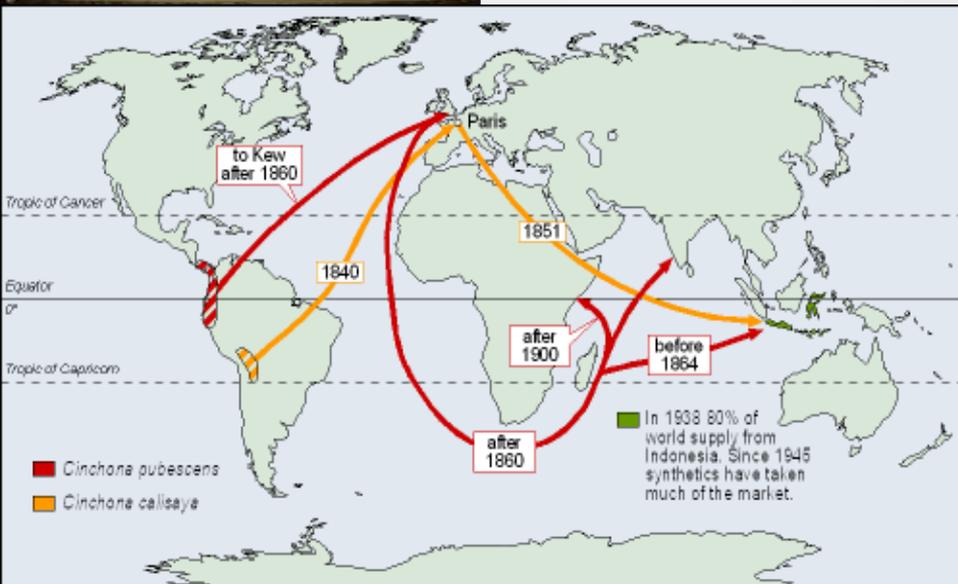
# Jesuits' powder

by Fr Eugene Nevins, SJ



College of Physicians, London, October 5, 1756.  
*We the Censors of the College, being informed that very large Quantities of a Bark imported from North America, have been lately sold by publick Sale at Garraway's Coffee House as Jesuits Bark, have procured Samples of it, and, calling to us the Master and Wardens of the Apothecaries Company, have examined it, and find, that though it has the Appearance, it has not the Taste or Quality of the true Jesuits or Peruvian Bark: We the Censors therefore, that we may not be wanting in an Affair of such Consequence to the Health and Lives of His Majesty's Subjects, think it our Duty to give this publick Notice to all Apothecaries, and other publick Dispensers of Medicines, desiring them not to purchase any Peruvian Bark without tasting and carefully examining it.*  
 Ambrose Dawson.  
 John Thomas Bott.  
 Antony Askew.  
 Nicolas Munchley.

London, October 12, 1756.



**A N A S T A S I S**  
**CORTICIS PERUVIÆ,**  
 SEU  
**CHINÆ CHINÆ**  
 DEFENSIO,  
**SEBASTIANI BADI GENVENSIS**  
*Præcipue variisque Nôbilitatibus olim Medicus,*  
 Et  
 Publicæ Sanitatis in Civitate Consulens.  
 Comæ  
**V E N T I L A T I O N E S**  
**IOANNIS JACOBI CHEFLETII,**  
 M. D. C. C. C. C. C. C. C.  
**VONSCI FORTVNATI PLEMPPI,**  
*Illustrissimi Medicorum.*  
*Opus in tres libros distinctum, & in six Dissertationes Medicas, & Philosophicas.*  
 ILLVSTRISSIMO D.  
**IOANNI LVCA DVRRATIO**

G E R V A , Typis Prael. Societatis "edendæ". M. DCC. LXXXI.  
 apud Joann. Valz.



THOMÆ SYDENHAM

Med. Doct.

METHODUS

# CURANDI FEBRES.

Propriis Observationibus  
Superstructa.

*Editio Secunda, priori multò auctior ac emendatior; cui etiam accessit Sectio Quinta de Peste sive Febre Pestilentiali.*

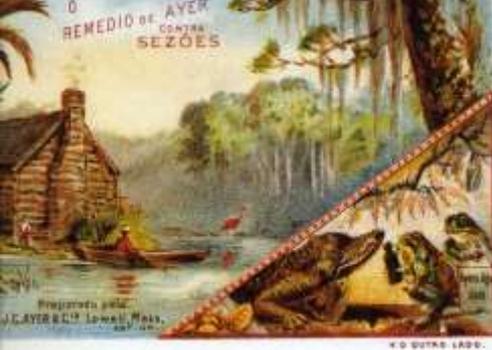
*Multa egerunt qui ante nos fuerunt, sed non peregerunt: multum adhuc restat opera, multumque restabit: neque ulli nato post mille secula præcidetur occasio aliquid adhuc adjiciendi. Seneca.*

LONDINI,

Impensis J. Crook, apud quem veniunt  
in Vico vocato Duck-lane propè  
Little Britain, MDCLXXVIII.



Thomas Sydenham (1624-1689) - 0



AGROTAT LIMB. CONIUX CHINCONIA FEBRIM  
CORTICE MIRABANO-POCULA TINCTA FUGANT

## EXPERIENCIAS CHYMICAS,

*Sobre a Quina do Rio de Janeiro comparada com outras.*

**P**ELA Secretaria de Estado dos Negocios da Guerra e da Marinha, foi commettida ao Exame da Academia Real das Sciencias de Lisboa a Analyse de huma porção de Casca amargosa, com o nome de Quina, que do Rio de Janeiro havia sido mandada ao Fysico Mór do Exercito; para que achando-se que continha os principios, e virtudes das Quinas officinaes, se fizesse uso della, principalmente nos Hospitales Militares.

Em consequencia, nomeou a Academia huma Commis-são, composta de quatro dos seus Membros, e pediu para os seus trabalhos o Laboratorio Chymico da Casa da Moeda, que immediatamente lhe foé franqueado, com ordem para que nelle se aprontasse tudo, quanto fosse necessario para aquella Analyse.

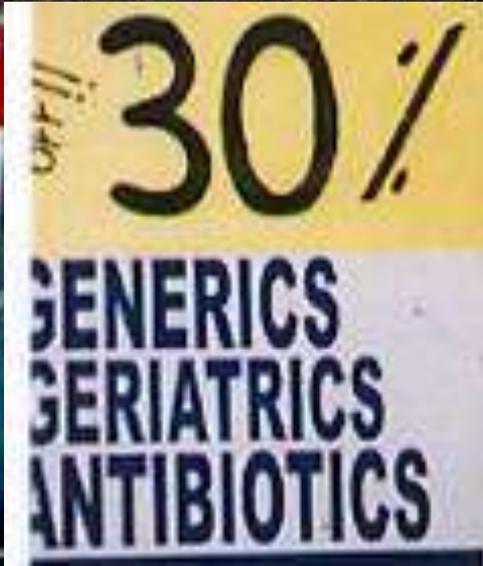
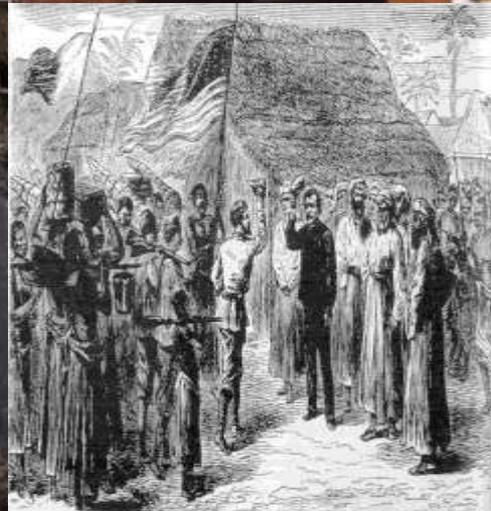
Este Laboratorio munido de todos os vasos e utencilios, que lhe são proprios; tinha contudo falta de muitos reagentes, principalmente d'aquelles que mais se alterão, os quacs seriamos obrigados a preparar; a não ser a franqueza do Sr. Alexandre Antonio Vandelli, que além de nos ajudar com o seu trabalho, nos forneceu os que nos forão necessarios, e ainda outros de que carecia o sobredito Estabelecimento.

Aos 17 de Junho, o primeiro dia em que ali concorremos, houve alguns pareceres sobre o Plano que deveriamos seguir nas nossas Experiencias. O interesse do objecto, a facilidade com que podiamos dispor dos meios que se nos offerociação; tudo nos convidava a fazer huma Analyse em toda a sua extenção: mas ao mesmo tempo, conhecendo que o nosso principal objecto era examinar aquella Casca relati-

VR.

“Experiências Chymicas, sobre a Quina do Rio de Janeiro Comparada com Outras”, escrita em 1811 por José Bonifácio, Sebastião Francisco de Mendo Trigo, João Croft e **Bernardino Antonio Gomes**, com a participação de Alexandre Vandelli”

(REVISTA TRIPLOV  
de Artes, Religiões e Ciências  
Nova Série | 2010 | [Número 05](#))





M A L A R I A  
**KILLS  
3000  
CHILDREN  
EVERYDAY**  
IN AFRICA. PLEASE HELP.  
[HTTP://MALARIANOWGISE.ORG](http://MALARIANOWGISE.ORG)



**AGAINST  
MALARIA  
FOUNDATION**

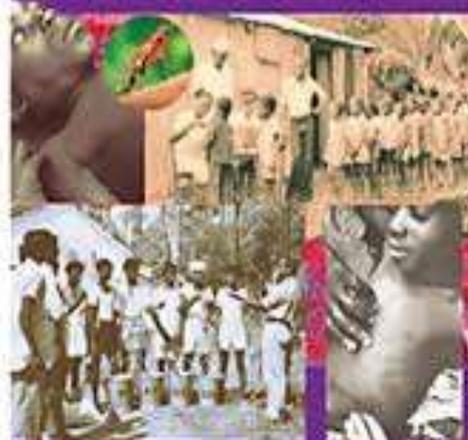
iea

Richard Toes & Roger Barr  
Founding Directors of IAEA

Malaria and  
the DDT Story



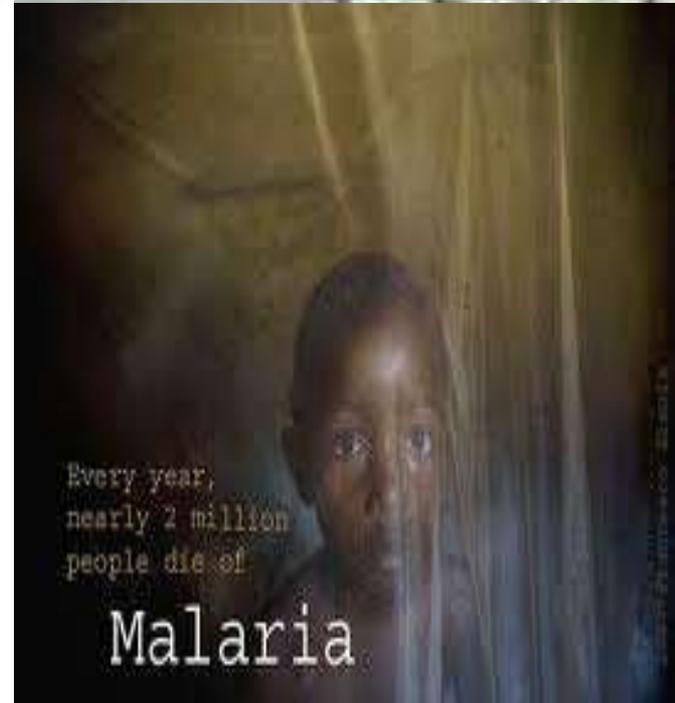
**HUMANITY'S BURDEN**  
A GLOBAL HISTORY OF MALARIA



JAMES L. A. WEBB JR.

Every year,  
nearly 2 million  
people die of

Malaria



# History of Malaria Control

## Time Line For Malaria Control

### Ancient Times

Early man attributed the fevers to evil spirits, angered deities, demons, or the black magic of sorcerers

Several thousand years ago

Babylonian cuneiform script attributes malaria to a god, pictured as a mosquito-like insect

800 BC

Indian sage Dhanvantari wrote that bites of mosquitoes could cause diseases, fever, shivering etc.

400 BC

Hippocrates attributed malaria to ingestion of stagnant water; also related the fever to the time of the year and to where the patients lived.

100 BC

Susruta Samhita in India associates fevers with the bites of the insects

### Early Civilizations

Use of mosquito nets recorded; Cleopatra reportedly used them

### Early Civilizations

Ancient Romans practiced drainage of swamps to prevent malaria

11 Century AD

Rice plantation prohibited near human dwellings

1717 AD

Lancisi linked malaria with poisonous vapours of swamps or stagnant water on the ground and revived the old idea that mosquitoes might play a role; proposed the draining of marshes to eradicate malaria

Late 1800

Malaria declines in the United States and Europe due mainly to draining swamps and removing mill ponds; better housing and better sanitation

1882 AD

Albert Freeman Africanus King, a US Physician, proposed to encircle the city with a wire screen as high as the Washington

Monument as a method to eradicate malaria from Washington, DC.

August 20, 1897

Ronald Ross demonstrates oocysts in mosquito gut, proving the role of mosquito in malaria transmission

1899

Ross attempted to eradicate malaria from England by forming 'mosquito brigades' to eliminate mosquito larvae from stagnant pools and marshes; organised a sanitation drive at Freetown, Sierra Leone with limited success

1901

William Gorgas of the US Army leads a successful anti mosquito drive in Havana, Cuba; pyrethrum used for the first time

1902

Ronald Ross invited to Ismailia by the Suez Canal Company to control malaria, which he does successfully.

1905-1910

Malaria controlled in Panama Canal site under the leadership of Ronald Ross and William Gorgas

1920

Paris Green first used in malaria control

1921-22

Larvivorous fish *Gambusia affinis* or mosquitofish used in the control of mosquitoes in California

1939

Malaria control drive conducted in Brazil under the leadership of Fred Sopper with great success

1955

The Global Malaria Eradication Programme launched by WHO with emphasis on vector control with DDT residual spraying

1965

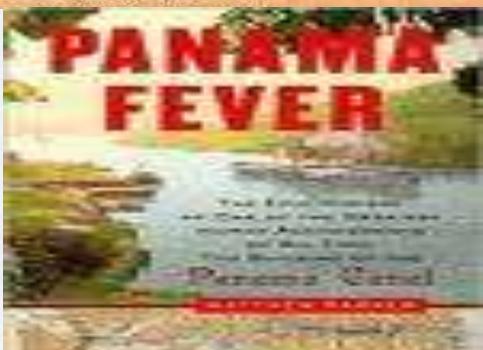
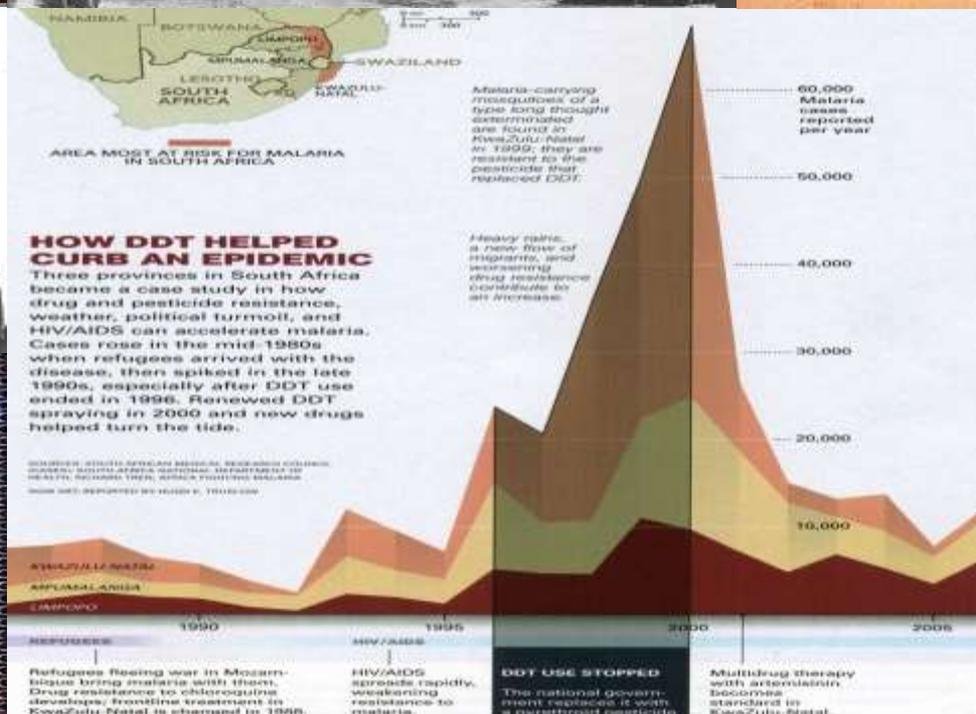
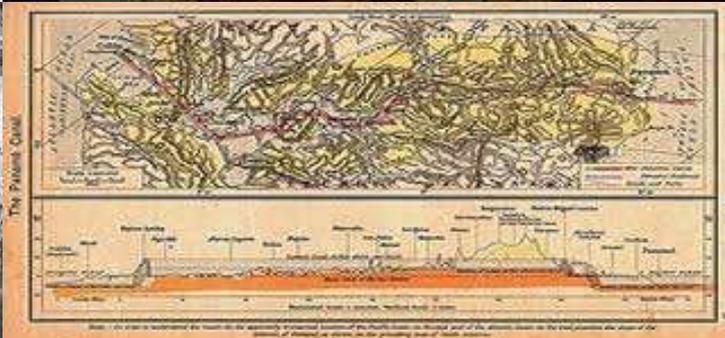
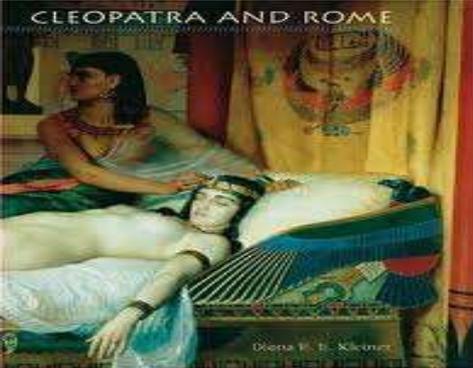
The Global Malaria Eradication Programme proved successful in Europe, but cases re-emerge in Asia

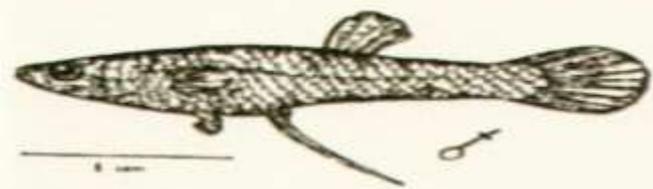
1969

The Global Malaria Eradication Programme abandoned

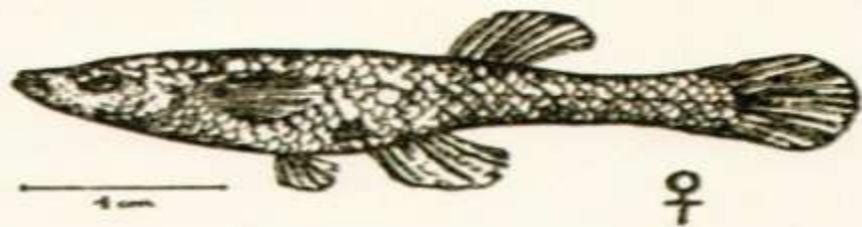
1998

Roll Back Malaria Campaign launched

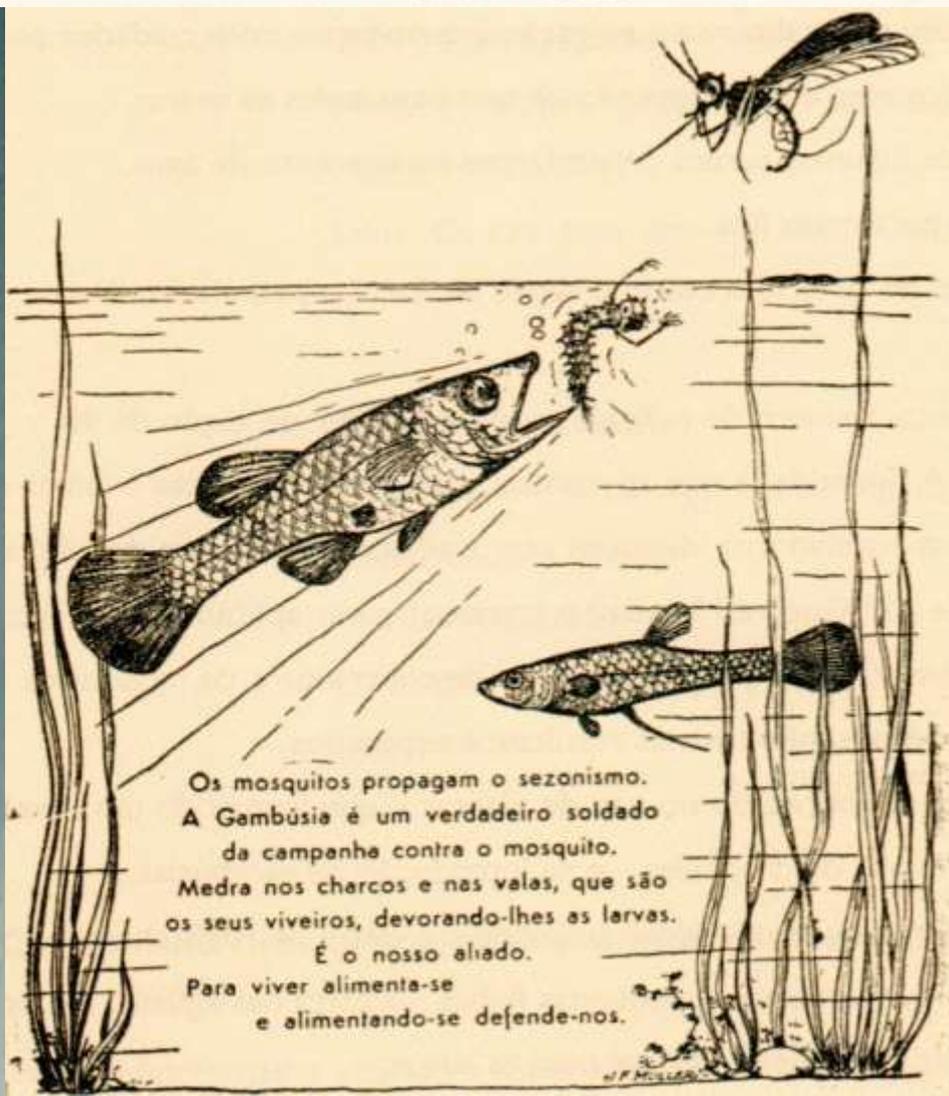




Gambouse mâle  
Gr. nat.



Gambouse femelle  
Gr. nat.



Mosquitoes at Malania

Drawn by Eugène Delacroix  
1848



(From a Mauritian Newspaper, 1908.)

On His Majesty's Service.

RR (cc) has much pleasure in accepting the kind invitation of RR

for the gathering at the Savoy on 10<sup>th</sup> October 1918.

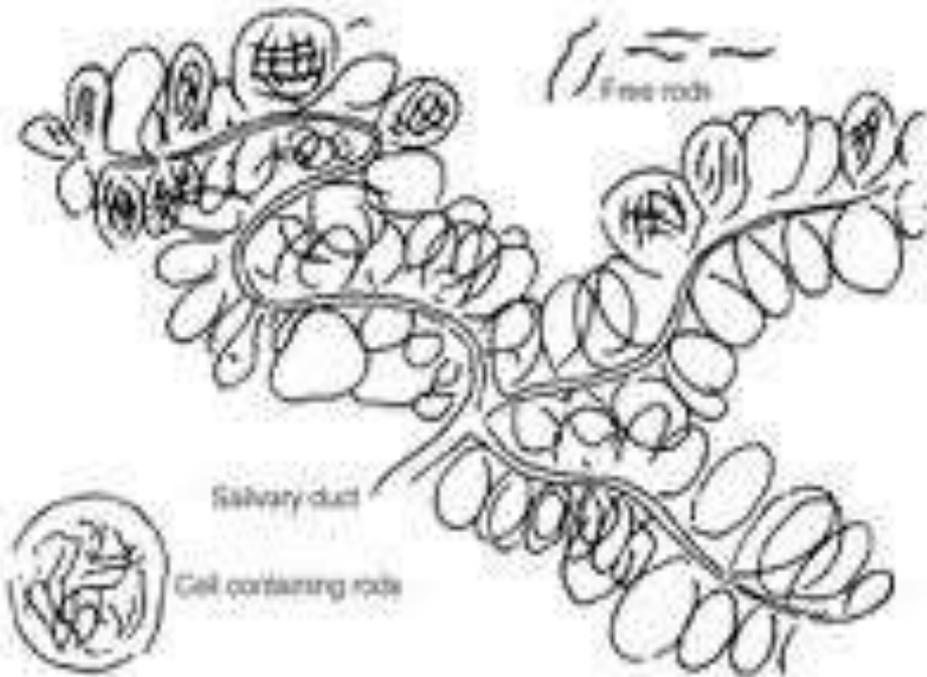
NOTICE  
Having to give  
service being  
given in  
OUR SERVICE  
ACCURATE  
PUNCTUAL  
AND DASHING  
AND NOT  
QUARANTEED  
BY THE FIRM

General Dispensing Board,  
Whitehall  
(18)

PARICATURES OF R. ROSS.



Figure 2. Alphonse Laveran's fight against malaria. Cartoon by B Moloch, published in *Chanteclair* (1908)



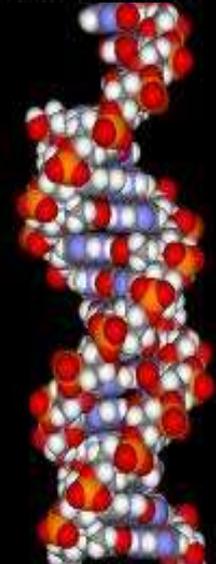
# First Malaria-Proof Mosquito: Genetic Manipulation Renders Them Completely Immune to the Parasite



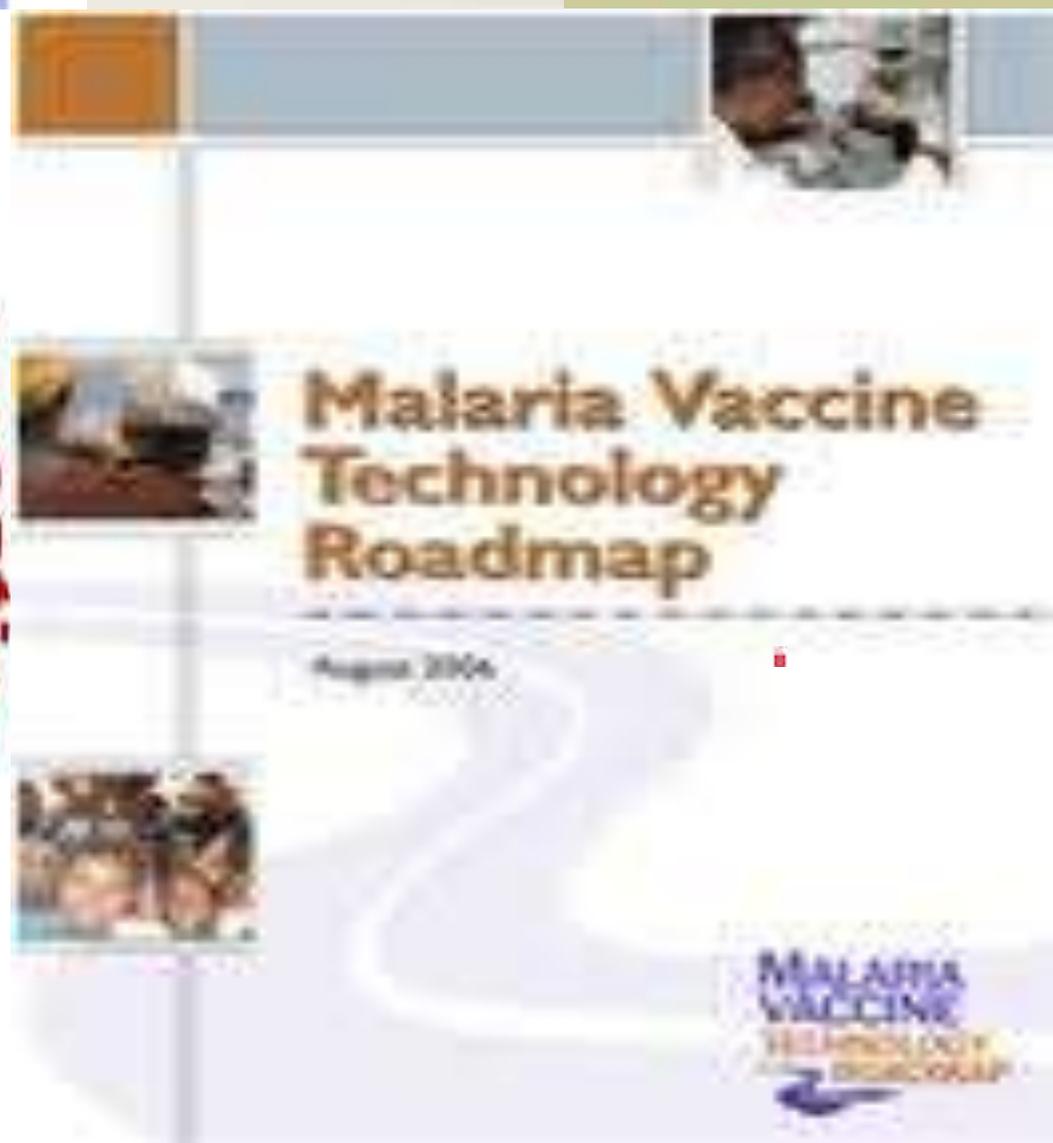
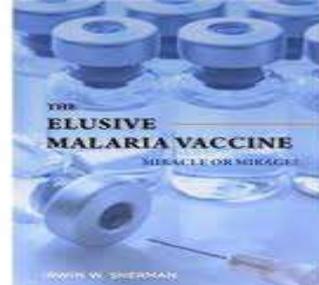
Genetically Modified Mosquito May End Malaria

Jul 16, 2010 9:05 AM CDT

Your source for the latest research news



## Genetics Modified Insects





*Plasmodium malariae* (Laveran, 1881)

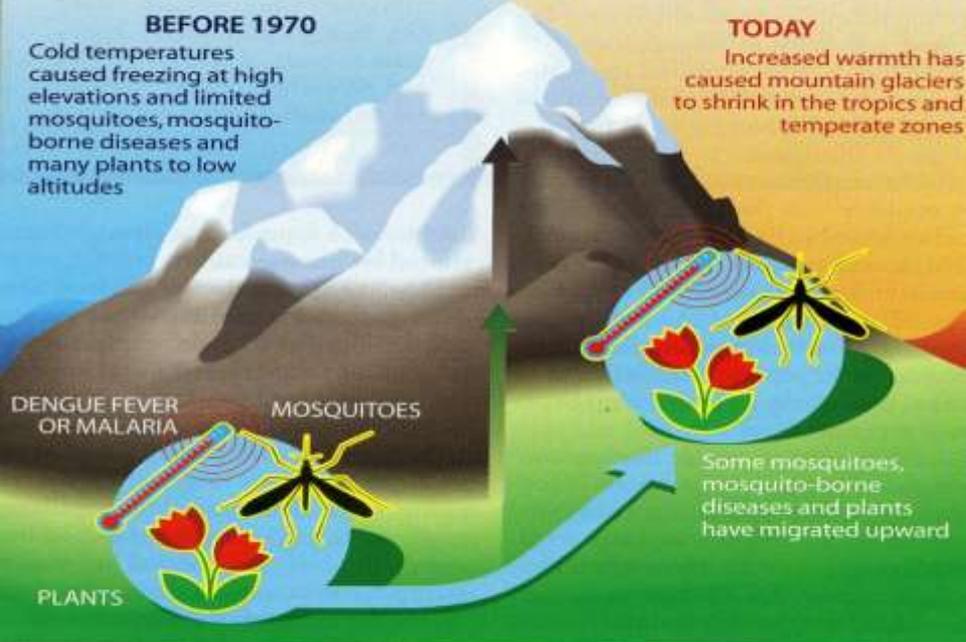


MALARIA... WEST NILE VIRUS... HANTAVIRUS... DENGUE... CHOLERA...

# Global Warming: The Hidden Health Risk

## Changes Are Already Under Way

Computer models have predicted that global warming would produce several changes in the highlands: summit glaciers (like North Polar sea ice) would begin to melt, and plants, mosquitoes and mosquito-borne diseases would migrate upward into regions formerly too cold for them (*diagram*). All these predictions are coming true. This convergence strongly suggests that the upward expansion of mosquitoes and mosquito-borne diseases documented in the past 15 years (*list at bottom*) has stemmed, at least in part, from rising temperatures.

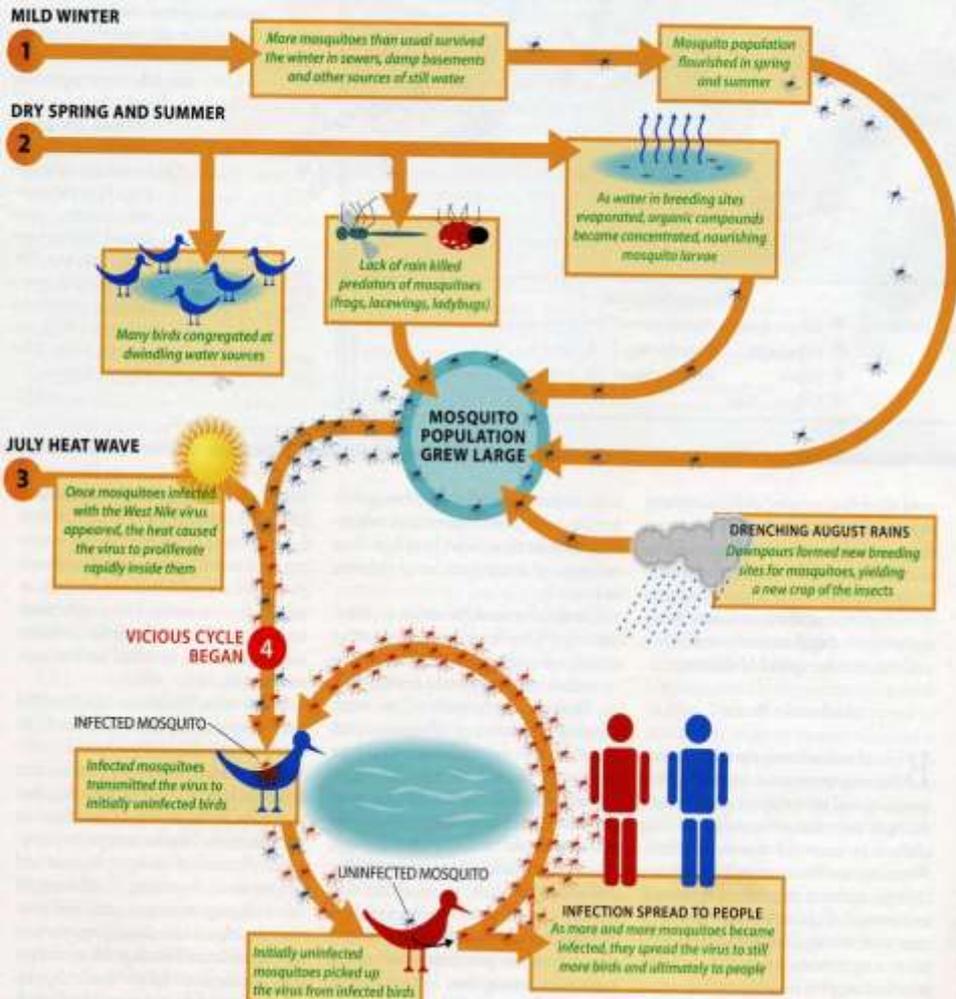


### WHERE DISEASES OR THEIR CARRIERS HAVE REACHED HIGHER ELEVATIONS

- |  |  |   |
|--|--|---|
| <b>Malaria</b><br>Highlands of Ethiopia, Rwanda, Uganda and Zimbabwe<br>Usamabara Mountains, Tanzania<br>Highlands of Papua New Guinea and West Papua (Irian Jaya) | <b>Dengue fever</b><br>San Jose, Costa Rica<br>Taxco, Mexico | <b>Aedes aegypti mosquitoes</b><br>(can spread dengue fever and yellow fever)<br>Eastern Andes Mountains, Colombia<br>Northern highlands of India |
|--|--|---|

## Weather and the West Nile Virus

This diagram offers a possible explanation for how a warming trend and sequential weather extremes helped the West Nile virus to establish itself in the New York City area in 1999. Whether the virus entered the U.S. via mosquitoes, birds or people is unknown. But once it arrived, interactions between mosquitoes and birds amplified its proliferation.



## Global warming and malaria: knowing the horse before hitching the cart

Paul Reiter

Address: Insects and Infectious Disease Unit, Institut Pasteur, 25-28 rue du Dr Roux, 75724 Paris, France

Email: Paul Reiter - paul.reiter@pasteur.fr

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Table 2. *Anopheles* species recorded from Europe, their status as malaria vector and geographic distribution.

Species	Status as malaria vector	Geographic distribution
<i>Anopheles algeriensis</i> Theobald	+	Mediterranean countries
<i>Anopheles claviger</i> (Meigen)	+	all of Europe
<i>Anopheles hyrcanus</i> (Pallas)	??	along Mediterranean coast
<i>Anopheles maculipennis</i> complex:		
<i>An. maculipennis</i> s.s. Meigen	??	temperate regions of Europe
<i>An. atroparvus</i> van Thiel	++	temperate regions of Europe, including Iberian peninsula
<i>An. beklemishevi</i> Stegnii & Kabanova	??	Sweden, Finland, Russia (Siberia)
<i>An. daciae</i> Linton, Nicolescu & Harbach	??	Romania, UK, other countries??
<i>An. labranchiae</i> Falleroni	+++	Southern Italy, Sardinia, Sicily, Corsica, Dalmatian coast
<i>An. melanoon</i> Hackett	??	Mediterranean coast
<i>An. messeae</i> Falleroni	??	all of Europe except Iberian peninsula, Southern Italy and Balkans/Greece
<i>An. sacharovi</i> Favre	+++	Southern Italy, Sardinia, Sicily, Corsica, Balkan, Greece, Turkey
<i>An. superpictus</i> Grassi	??	Corsica, Italy, Balkans, Greece
<i>Anopheles petragrani</i> del Vecchio	??	along Mediterranean coast
<i>Anopheles plumbeus</i> Stephens	??	temperate regions
<i>Anopheles sergentii</i> (Theobald)	+++	Sicily

+, ++, +++ Proven malaria vectors, in ascending importance; ?? Malaria vector status uncertain.

### Abstract

Speculations on the potential impact of climate change on human health frequently focus on malaria. Predictions are common that in the coming decades, tens – even hundreds – of millions more cases will occur in regions where the disease is already present, and that transmission will extend to higher latitudes and altitudes. Such predictions, sometimes supported by simple models, are persuasive because they are intuitive, but they sidestep factors that are key to the transmission and epidemiology of the disease: the ecology and behaviour of both humans and vectors, and the immunity of the human population. A holistic view of the natural history of the disease, in the context of these factors and in the precise setting where it is transmitted, is the only valid starting point for assessing the likely significance of future changes in climate.

## 2. Will malaria return to North-West Europe?

Willem Takken, Piet A. Kager and Jan Peter Verhave

### Abstract

Historically, malaria was widespread in Europe, reaching as far north as the United Kingdom, the Netherlands and even Sweden and Finland. In southern Europe, malaria was caused by *Plasmodium falciparum*, *P. vivax* and *P. malariae*, but further north only the latter two parasite species were found. The main vector species was *Anopheles atroparvus* Van Thiel, although *An. messeae* Falleroni was suspected to be a vector in Central Europe and Scandinavia. The disease was eradicated from Europe in the second half of the 20th century with DDT and modern drugs. In spite of a near-constant and large number of cases of imported malaria, as well as the proven presence of anopheline vectors, the return of local malaria transmission has not been observed, presumably because of a lack of a sufficiently dense cluster of parasite carriers and an efficient health care system that is able to diagnose and treat malaria patients successfully. It is considered whether climate change might alter this situation, allowing for higher mosquito densities and more favourable environmental conditions for *Plasmodium* development in the mosquito vectors. We argue that under climate change, causing higher temperatures and intensified precipitation, conditions for local malaria transmission will improve, but that the public health measures will preclude the building up of an infectious parasite reservoir. So people might be exposed to more mosquito bites, but the chance of such mosquitoes becoming infected with malaria parasites will not alter significantly compared to the current situation. Therefore, climate change is unlikely to affect the malaria-free state of North-West Europe.



Figure 1  
Malaria in Norway, 1880–1910 (37). Unpublished map by Lars Holden and Larry Holden (with permission of the authors).

RESEARCH

Open Access

## Malaria resurgence risk in southern Europe: climate assessment in an historically endemic area of rice fields at the Mediterranean shore of Spain

Sandra Sainz-Elipe<sup>1</sup>, Jose Manuel Latorre<sup>1</sup>, Raul Escosa<sup>2</sup>, Montserrat Masià<sup>2</sup>, Marius Vicent Fuentes<sup>1</sup>, Santiago Mas-Coma<sup>1</sup>, Maria Dolores Bargues<sup>1\*</sup>

### Abstract

**Background:** International travel and immigration have been related with an increase of imported malaria cases. This fact and climate change, prolonging the period favouring vector development, require an analysis of the malaria transmission resurgence risk in areas of southern Europe. Such a study is made for the first time in Spain. The Ebro Delta historically endemic area was selected due to its rice field landscape, the presence of only one vector, *Anopheles atroparvus*, with densities similar to those it presented when malaria was present, in a situation which pronouncedly differs from already assessed potential resurgence areas in other Mediterranean countries, such as France and Italy, where many different *Anopheles* species coexist and a different vector species dominates.

**Methods:** The transmission risk was assessed analysing: 1) climate diagrams including the minimum temperature for *Plasmodium falciparum* and *Plasmodium vivax* development; 2) monthly evolution of the Gradient Model Risk (GMR) index, specifying transmission risk period and number of potential *Plasmodium* generations; 3) ecological characteristics using remote sensing images with the Eurasia Land Cover characteristics database and the monthly evolution of the Normalized Difference Vegetation Index (NDVI); 4) evaluation of *A. atroparvus* population dynamics.

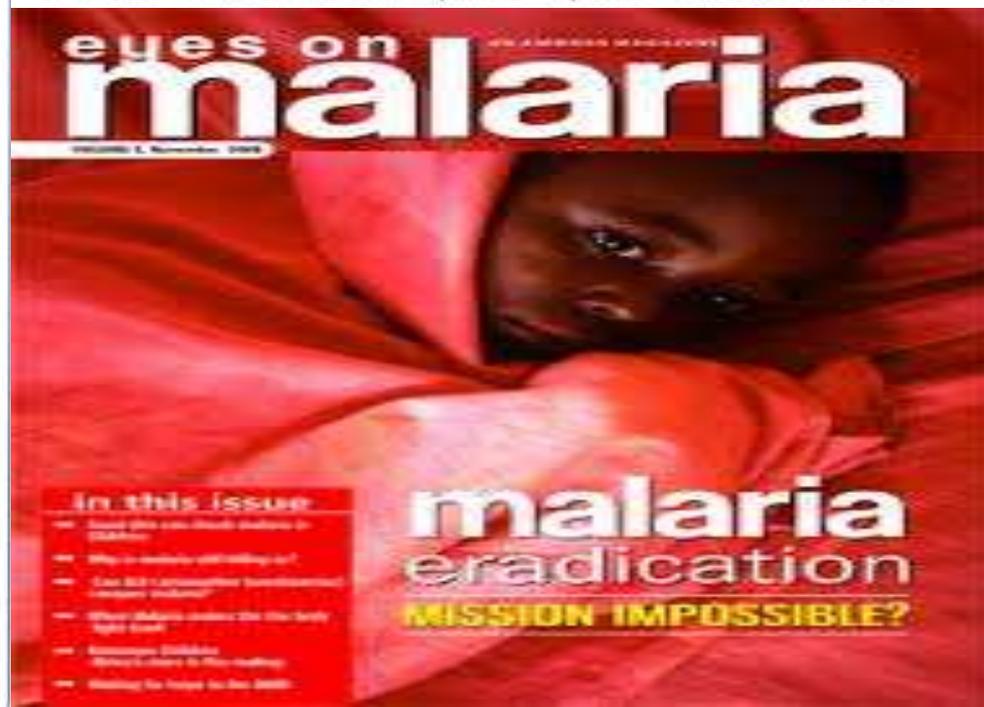
**Results:** Climatological analyses and GMR index show that a transmission risk presently exists, lasting from May until September for *P. falciparum*, and from May until October for *P. vivax*. The GMR index shows that the temperature increase does not actually mean a transmission risk increase if accompanied by a precipitation decrease reducing the number of parasite generations and transmission period. Nevertheless, this limitation is offset by the artificial flooding of the rice fields. Maximum NDVI values and *A. atroparvus* maximum abundance correspond to months with maximum growth of the rice fields.

**Conclusions:** The Ebro Delta presents the ecological characteristics that favour transmission. The temperature increase has favoured a widening of the monthly potential transmission window with respect to when malaria was endemic. The combined application of modified climate diagrams and GMR index, together with spatial characterization conforms a useful tool for assessing potential areas at risk of malaria resurgence. NDVI is a good marker when dealing with a rice field area.

## Airport malaria: a review

M. Isaäcson<sup>1</sup>

*Cases of malaria occasionally arise among individuals who have never visited a malarious area. Such patients, who also lack a history of blood transfusions or intravenous drug abuse, are usually shown to have "airport malaria". Most reports of airport malaria consist of case histories, although some epidemiological reviews have also appeared. The clinical and epidemiological features of 29 cases of airport malaria that were reported in Europe from 1969 to 1988 are reviewed here. Although airport malaria is rare, the apparent absence of risk factors for the disease in a patient's history can result in delays in diagnosis and appropriate treatment. Tests to exclude malaria should therefore be carried out on patients who work at or live near an international airport and who present with acute febrile illnesses.*



# Malaria, a Neglected Factor in the History of Greece and Rome

W. H. S. Jones



Εικόνα 1. Τόπος κατοικίας ασθενών με ελονοσία χωρίς ιστορικό ταξιδιού σε ενδημική χώρα. Ελλάδα, περίοδος 2011 (έως 16.09.2011)

Πίνακας 1. Τόπος κατοικίας ασθενών με ελονοσία χωρίς ιστορικό ταξιδιού σε ενδημική χώρα σε Έλληνες ασθενείς. Ελλάδα, περίοδος 2011 (έως 16.09.2011)



Περιφερειακή Ενότητα κατοικίας	Αριθμός ασθενών	Ασθενείς ανά 100.000 πληθυσμού*
Λακωνίας	10	8,61
Δήμος Ευρώτα	10	50,50
Ευβοίας	2	0,97
Βοιωτίας	1	0,80
Ανατολικής Αττικής	2	0,50
Λάρισας	1	0,35
<b>Σύνολο χώρας</b>	<b>16</b>	<b>0,14</b>

\*Υπολογίστηκε με βάση στοιχεία πληθυσμού από την Ελληνική Στατιστική Αρχή (εκτιμήσεις 2008).

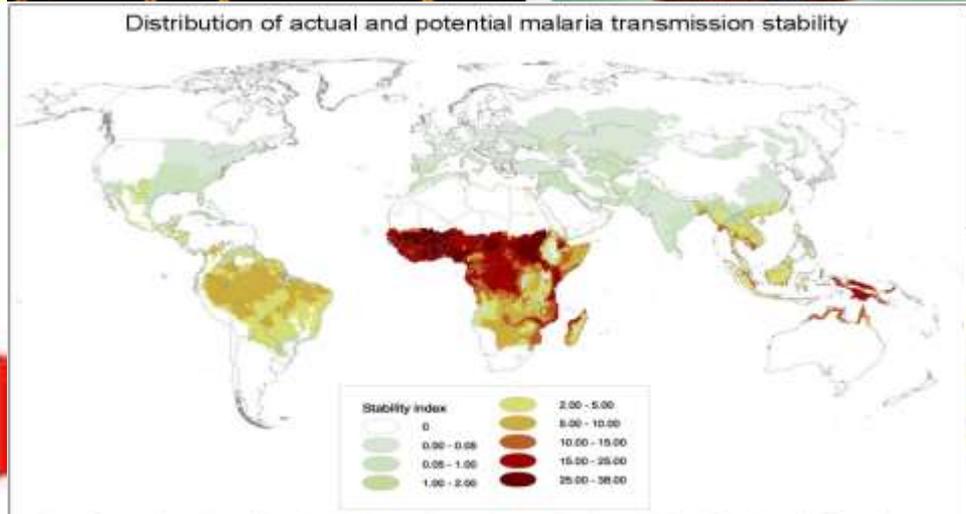
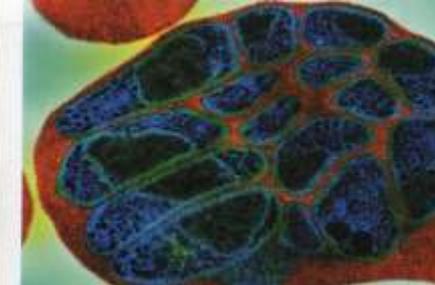
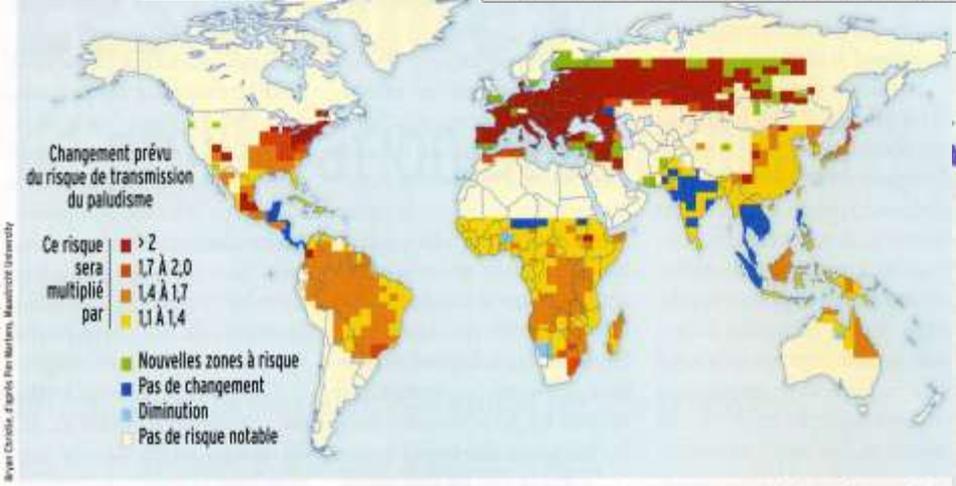


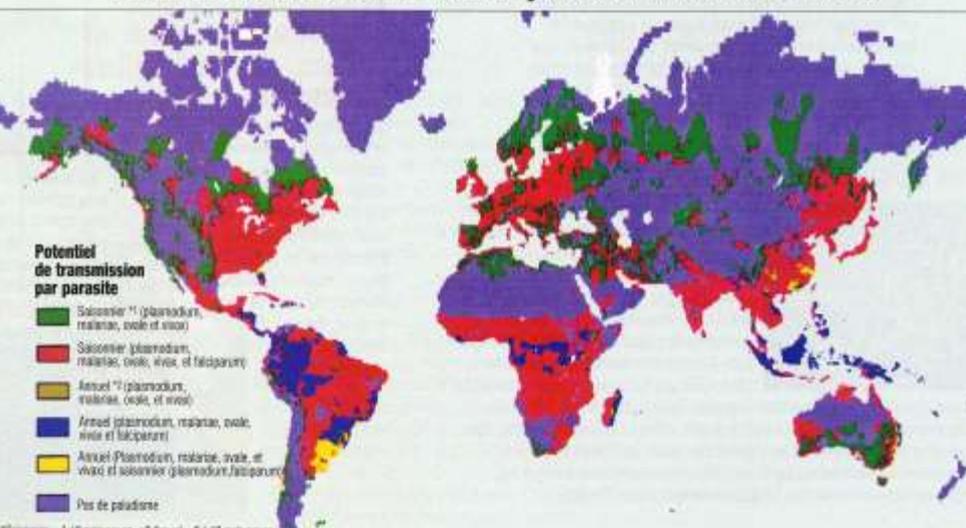
Figure 1 from Anthony Kiszewski, Andrew Mellinger, Andrew Spielman, Pia Malaney, Sonia Erlich Sachs, and Jeffrey Sachs. A Global Index Representing The Stability of Malaria Transmission. *Am J Trop Med Hyg* 2004 70:486-498.



2. LE RISQUE DE TRANSMISSION DU PALUDISME aux alentours de 2020 aura augmenté dans de nombreuses parties du monde (en comparaison avec le risque moyen dans les années 1961 à 1990), selon des

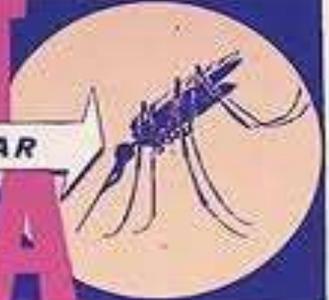
projections fondées sur un accroissement de la température de 1 °C. L'analyse ne tient compte que de la variation de température et pas des autres facteurs qui agirait sur l'extension du paludisme.

### Potentiel de transmission du paludisme en 2025-2045



**Crusade against Malaria**  
Malaria-ipca.com, an exhaustive online resource centre on malaria

**PREVENT MALARIA**  
SHORTEN THE WAR



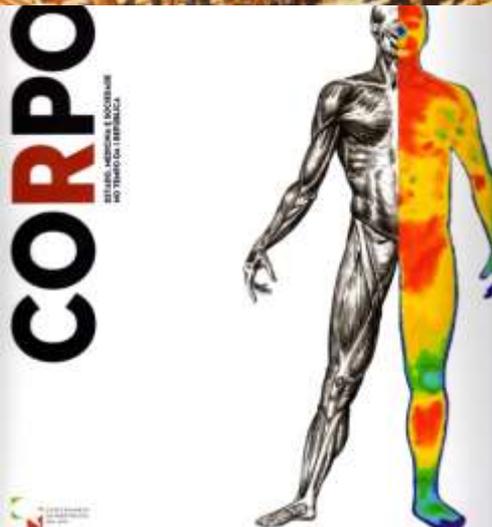
# A MALÁRIA EM PORTUGAL CONTINENTAL

**malaria**  
**NO MORE**

**Malaria Menace!!!**  
People who have never had a malaria infection such as young children, travelers and pregnant women are more likely to have severe symptoms from malaria.



**Read More**



**INSTITUTO PASTEUR DE LISBOA**

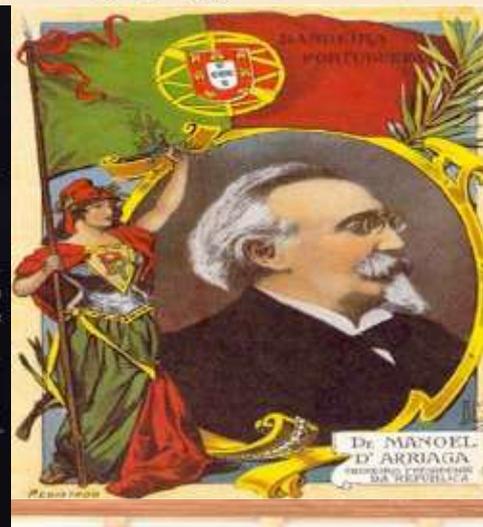
*seros vacinas análises clínicas*

**N.º DE ÓBITOS ANUAIS\***  
(MÉDIA DE 1916 A 1925)

TUBERCULOSE	8980	PNEUMONIA	3849	CÂNCER	1777
MEMINGITE	1734	VARÍOLA	1705	FEBRE TIFOIDE	1163
SARDANHO	819	TOSSE CONVULSA	703	CIROSE DO FÍGADO	671
DIFTERIA	563	TIFO EXANTEMÁTICO	334	SEZONISMO	229
ESCARLATINA	30				

**PIQUENOS DA RAIVA!**

**destruir o rato e matar o UFA**



**RAIVA**  
3153 PESSOAS TRATADAS  
1918

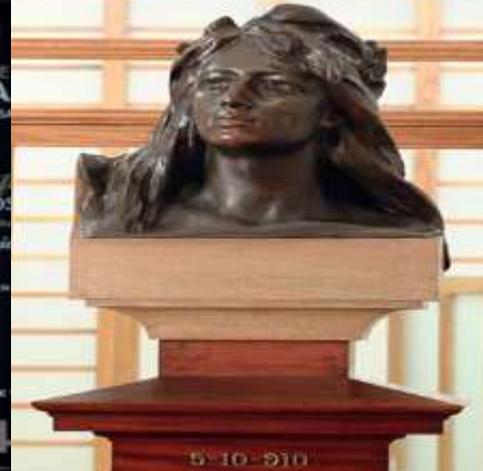
**1892** INSTITUTO CÂMARA PESTANA

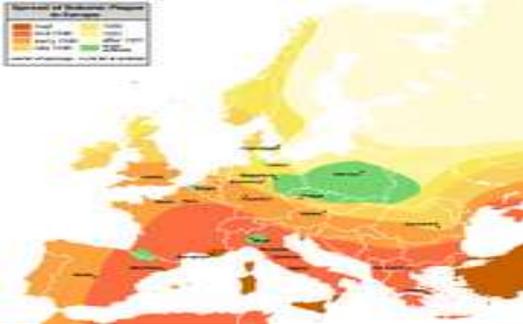
**PREPARAÇÃO DE SOROS TERAPÉUTICOS**  
**498 409**  
Total de soros de s. e. e. preparados de 1911 a 1938

**MORTALIDADE DOENÇA INFECTO-CONTAGIOSA**

*fame, epidemia*  
**TUBERCULOSE**  
mas, catenra, gripe pneumónica

**TOHO DE MORTALIDADE**  
em 1918 e 1919  
**26,4**





# Ricardo Jorge



## SOBRE A SAÚDE PÚBLICA ...

“... o formigueiro humano germina em si próprio o veneno das sua destruição ... há um estendal de vergonhas, de males e incúrias ... é forçoso lavrar um protesto contra tanto desleixo, contra tanta inépcia, contra tanta loucura criminosa ... não são só as revoadas de contágio que desafiam a debelação, mas também as endemias tenazes ... tal como o sezonismo que estende pelo País largas manchas ... há mais de 20 anos que as estâncias de saúde se preocupam com o estudo e combate do flagelo sem colher a realização dos seus propósitos ... hoje basta dizer que somos o único país da Europa sem guerra organizada contra o sezonismo ... esta é, como poucas, uma questão patriótica ... não será temeridade afirmar-se que a infecção sazonal avulta entre as piores pragas colectivas que flagelam a população portuguesa ... contra o flagelo da tuberculose inaugurou-se, mas não se activou o combate ... contra as moléstias venéreas, não há um único serviço de dispensário montado, como hoje o mandam os mais elementares princípios ...” (sic.)

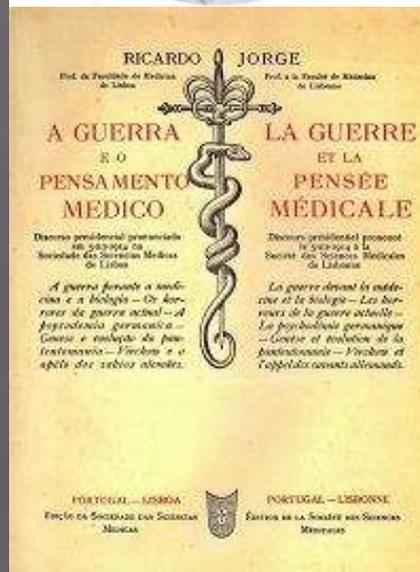


Figura 3 – O Professor Ricardo Jorge e a Peste Bubónica no Porto. Caricatura de Manuel Monterroso (1876-1968).

Percursos da Saúde Pública nos séculos XIX e XX - a propósito de Ricardo Jorge



Imagem: Arquivo de Ricardo Jorge, Porto, Arquivo de Ricardo Jorge, Porto, Arquivo de Ricardo Jorge, Porto



1093

1902-1060

1073

1900

N.º 5

N.º 20

PROPHYLAXIA INDIVIDUAL  
DO  
**PALUDISMO**  
(BREVE ESTUDO)

DISSERTAÇÃO INAUGURAL  
APRESENTADA À  
ESCOLA MEDICO-CIRURGICA DO PORTO  
POR  
*Sebastião Eduardo Cezar de Sá*

111/4 ENC

PORTO  
Officina de «Comércio do Porto»  
108—Rua de «Comércio do Porto»—111  
—  
1902

BREVE ESTUDO  
SOMAS O  
**PALUDISMO**

(ACCESOS PERNICIOSOS EM HOSPITAL)  
DISSERTAÇÃO INAUGURAL  
APRESENTADA À  
ESCOLA MEDICO-CIRURGICA DO PORTO  
POR  
*ANTONIO MACHADO ACABADO*

PORTO  
TYP. A VAGOR DA IMPRENSA LITTERARIA E TYPOGRAPHICA  
27, rua de S. João, 124  
1900

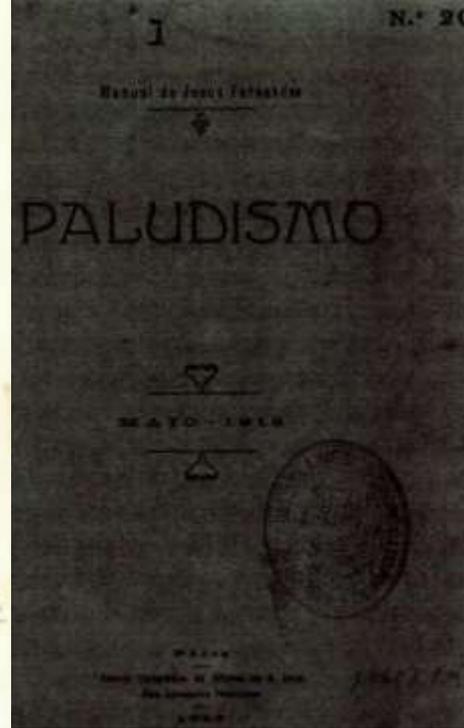
112/3 ENC

BREVE ESTUDO  
SOBRE  
ETIOLOGIA E PROPHYLAXIA  
DO  
**PALUDISMO**

DISSERTAÇÃO INAUGURAL  
APRESENTADA À  
ESCOLA MEDICO-CIRURGICA DO PORTO  
POR  
*Antonio Loureiro Dias*

105/6 ENC

PORTO  
TYP. DE A. F. YACONTELLER, SUCCESOR DE  
21, Rua de S. Vicente, 28  
1901



03-10

: Contribuição  
para o estudo  
do paludismo  
larvado : : :

Tese de doutoramento apresentada  
à Faculdade de Medicina da Universi-  
dade do Porto, no ano de 1923  
por  
**JOÃO PEREIRA RIBEIRO**

203110 ENC

IMPRENSA E BARRAGEM SA  
CASA TYPOGRAPHICA  
COIMBRA - 1923

VIII N.º 88  
FACULDADE DE MEDICINA DO PORTO

*Arnando Rodrigues*  
**A PATOGENIA DA FEBRE BILIOSA  
HEMOGLOBINÚRICA**

TESE DE DOUTORAMENTO  
1921 apresentada à FACULDADE  
Faculdade de Medicina do Porto

MARÇO DE 1921

11518 ENC

PORTO  
Tip. a vapor da "Enciclopedia Portuguesa"  
47, Rua Cândido das Rios, 48

1172

J. CONDILLAC PINTO  
*1074*  
ESTUDO  
SOBRE A ETIOLOGIA E A PROPHYLAXIA  
DA

**MALARIA**

DISSERTAÇÃO INAUGURAL  
APRESENTADA À  
ESCOLA MEDICO-CIRURGICA DO PORTO

PORTO  
Officina de «Comércio do Porto»  
108—Rua de «Comércio do Porto»—111  
—  
1901

1904

*A. B.*  
PROPHYLAXIA  
DA  
**INFECCÃO PALUSTRE**

Dissertação inaugural para acto grande,  
seguida de nove proposições,  
e apresentada à Faculta. Medico-Cirurgica do Porto  
PARA OSES DEFFREYRA  
POR  
**MATHEUS AUGUSTO RIBEIRO DE SAMPAIO**

NOA A PROPHYLAXIA DO CEE.º RES.  
**ILLIDIO ATRÉS FERREIRA DO VALLE**  
LEITOR DE QUARTA CARRERA

PORTO  
Imprensa Portuguesa  
—  
1912

111/4 ENC

Introdução ao Estudo

DAS

# FEBRES DE GÔA

(Uma pagina de Pathologia Colonial)

DISSERTAÇÃO INAUGURAL  
APRESENTADA A

ESCOLA MEDICO-CIRURGICA DO PORTO



144/2 EHC

PORTO

Typ. a vapor de "Enciclopedia Portugueza Illustrada,"  
Rua N. S. do Anjo, 47 e 49

1910

Antonio de Mattos Pinto d'Azevedo

ob. 2.

## PROPHYLAXIA

DAS

# Doenças endemicas dos paizes quentes

Dissertação inaugural

apresentada á

Escola Medico-Cirurgica do Porto



PORTO

Typ. a vapor de "Enciclopedia Portugueza Illustrada,"  
11, Rua do Rocio D. Anjo, 47

1907

132/2 EHC

## BREVE ESTUDO

SOBRE A

# Endemo-epidemia palustre

DISSERTAÇÃO INAUGURAL  
APRESENTADA Á

Faculdade de Medicina do Porto

JULHO DE 1912

Tip. a vapor de PAP. E TYP. ALMEIDA  
de ALMEIDA & FILHOS - Largo do Lopo, 16  
1912 - 1913

José de Palma Gomes

ob. 13

# O Paludismo e o Arrhenal

(ALGUMAS OBSERVAÇÕES)

Dissertação inaugural  
apresentada á  
Escola Medico-Cirurgica do Porto



PORTO  
Cooperativa Graphica  
R. Empedado, 21  
1900

130/12 EHC

N.º 10.

# CHOROGRAPHIA PALUSTRE

DE

# PORTUGAL

DISSERTAÇÃO INAUGURAL

APRESENTADA Á

ESCOLA MEDICO-CIRURGICA DO PORTO



PORTO

TYP. A VAPOR DA REAL OFFICINA DE S. JOSÉ  
Rua Alexandre Herculano

1899

# Paludismo em Portugal: Breve Resenha Histórica I

- ▣ 1844: Escola Médica de Goa
- ▣ 1859: Comissão Nacional para analisar a relação entre a orizicultura e a Malária
- ▣ 1860: Publicação do Relatório da Comissão
- ▣ 1881: Inauguração do Laboratório Municipal de Higiene de Lisboa
- ▣ 1882: Inauguração do Gabinete de Microbiologia de Coimbra e Laboratório Municipal de Higiene do Porto
- ▣ 1887: Ensino da Medicina Tropical na Escola Naval
- ▣ 1888: 1º Artigo publicado por José Joyce “Das Formas Larvares do Impaludismo” (Medicina Contemporânea, Hospital da Marinha)
- ▣ 1892: Criados os Institutos Bacteriológicos de Lisboa e Porto
- ▣ 1895: Inauguração da Escola de Medicina Tropical (futuro IHMT)
- ▣ 1900: Inauguração do Instituto Câmara Pestana e realização do 1º Inquérito Nacional do Impaludismo
- ▣ 1901: Ricardo Jorge elabora o “Regulamento Geral dos Serviços de Saúde e Beneficência Pública” e cria-se o Instituto Central de Higiene (futuro INSA); Nomeada uma Comissão para Estudo e Combate ao Paludismo cujas recomendações são discutidas na Sociedade de Ciências de Lisboa
- ▣ 1901: 1ª Missão de Combate á Tripanosomose em Angola
- ▣ 1904: Primeiro estudo sobre o Anopheles em Portugal
- ▣ 1906: Realização em Lisboa do Congresso Internacional de Medicina
- ▣ 1911- Criação da Comissão Nacional de Defesa Anti-Sezonática
- ▣ 1920 - 1942: Intensificação da orizicultura em Portugal e Missão de Combate e Estudo da Malária na Aldeia Galega (Montijo)
- ▣ 1926: Reorganização geral dos Serviços Nacionais de Saúde Pública
- ▣ 1929: Assistentes do Instituto Câmara Pestana frequentam Curso Internacional de Malariologia organizado pela Sociedade das Nações; Início do combate ao Sezonismo, e Início do Ensino das Visitadoras Sanitárias

# Paludismo em Portugal: Breve Resenha Histórica II

- 1930 Abertura da Estação Experimental de combate ao Sezonismo em Benavente
- 1932: Abre a 1ª Estação Anti-Sezonática (A. Sal) e estabelece-se o 1º contacto entre o Governo Português e a Fundação Rockefeller
- 1933: Chegada dos 1ºs Estagiários Portugueses à John's Hopkins School of Hygiene e realização do 1º Inquérito Nacional, publicação do 1º Relatório Nacional (F. Cambournac), e Organização dos Serviços Anti-Sezonáticos
- 1934: Constituída a Comissão Reguladora do Comércio de Arroz e início das actividades da Estação Anti-Sezonática de A. Moura (futuro Instituto de Malariologia, com fins de Combate, Investigação e Formação de Técnicos); Início da taxaçoão do Comércio do Arroz c/ finalidade de subsidiar o Combate ao Sezonismo; Criação dos Centros de Saúde
- 1937: Início da actividade das Ambulâncias - Laboratórios
- 1943: Início da publicação dos Anais do IHMT
- 1939: O Sezonismo é considerado DDO, e F. Cambornac é nomeado Director do Instituto de A. Moura, altura em que este é doado pela F. Rockefeller ao Estado Português
- 1940: Introdução infrutífera de Estírpes de Mosquitos incapases de transmitir Plasmodium em Canha, e início da utilização do DDT
- 1941: Inauguração de uma pequena Enfermaria no I. de A. Moura
- 1946: F. Cambournac representa Portugal na Reunião inaugural da OMS
- 1953: Relatório sobre a Malária em S. Tomé e Príncipe
- 1954: Realização dos 1ºs Cursos Internacionais de Malariologia da iniciativa da OMS e dirigidos por F. Cambournac
- 1958: Malária endémica é erradicada de Portugal Continental
- 1959: Celebração do Acordo de Coordenação Anti-Palúdica Ibérico
- 1971: O Instituto de Malariologia é integrado no INSA
- 1973: O Instituto de A. Moura passa a dedicar-se ao estudo das Zoonoses e a Europa é declarada Região não Endémica de Malária pela OMS
- 1975: 1 caso de transmissão autóctone descrito no Distrito de Beja
- 1993: O Instituto de A. Moura passa a designar-se CEVDI



# MINISTÉRIO DO INTERIOR

Direcção Geral de Saúde

Repartição de Saúde

Parque Sanitário



Comissão Reguladora do Comércio do Arroz → DIRECÇÃO DE SERVIÇOS ANTI-SEZONÁTICOS ←

Fundação Rockefeller

INSTITUTO DE MALARIOLOGIA

Secção Administrativa

Secção de Publicidade e Propaganda

Secção de Engenharia

Secção Científica

Secretaria  
Expediente e Arquivo

Contabilidade

Armazém - depósito

Transportes

Biblioteca

Boletim

Estatística

Cartazes,  
Fotografia

Bonifica  
Hidráulica

Habitacões  
e  
Saneamento

Protecção  
Mecânica

Ensino

Investigação



## SECÇÃO TÉCNICA DE LUTA ANTI-SEZONÁTICA

ESTAÇÕES

POSTOS



Montemor-O-Velho

Benavente

Idanha-A-Nova

Alcácer do Sal

Pocinho

Soure

Ponte de Sôr

Azambuja

Dispensários:  
(1)  
Oliveira do Bairro  
Louriçal

Consultas  
Ambulantes:  
Tentugal  
Arazede  
Vila Verde  
Fôja  
Rol  
S. Silvestre

Dispensários:  
Coruche  
Vendas Novas  
Aguas de Moura

Consultas  
Ambulantes:  
Samora Correia  
Santo Estevão  
Salvatera de Magos  
Marinhais  
Muge

Dispensários:  
Monfortinho

Consultas  
Ambulantes:  
Ladoeiro  
Medelim  
Monsanto  
Proença-A-Velha  
Zebreira

Dispensários:  
Comporta  
Aljezur  
Quarteira

Consultas  
Ambulantes:  
Santa Suzana  
Grândola  
D. Rodrigo  
Torão  
Palma  
Parchanas

Consultas  
Ambulantes:  
Muxagata  
Touga  
Freixo de Numão  
Junqueira  
Horta da Vilarça

Consultas  
Ambulantes:  
Pombal  
Vila Nova de Anços  
Sebal Grande  
Ameal

Consultas  
Ambulantes:  
Galveias  
Montargil  
Bemposta  
Longomel

Consultas  
Ambulantes:  
Carregado  
Vila Franca de Xira  
Cartaxo  
Valada  
Vale de Santarém  
Ota

(1) **Dispensário:** Chamava-se dispensário ao destacamento duma visitadora sanitária ou dum auxiliar de laboratório para uma localidade onde actua, sob a direcção da Estação respectiva.



**INSTITUTO DE MALARIOLOGIA  
ÁGUAS DE MOURA — PORTUGAL**

Temos o prazer de anunciar que, com a autorização de Sua Excelência o Ministro do Interior, e da Direcção Geral de Saúde Pública, o Doutor Francisco José C. Cambournac foi nomeado Director do Instituto de Malaria, e tomará posse do novo cargo no dia 2 do próximo mês de Dezembro.

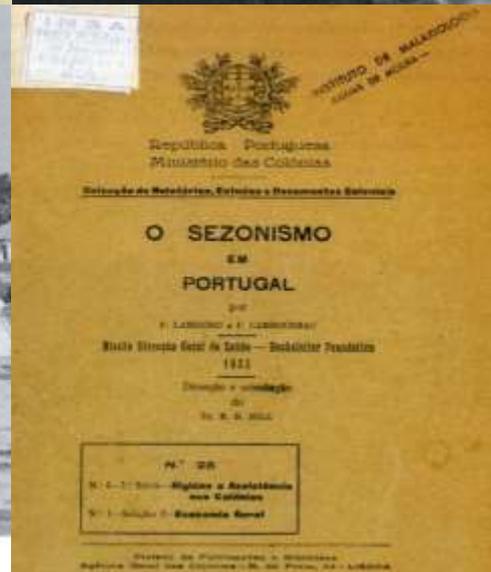
We are pleased to announce that Doctor Francisco José C. Cambournac has been appointed Director of the Malaria Institute, as of December 2, 1939.

ÁGUAS DE MOURA  
November 15, 1939.

*Dr. Pedro B. Hill*  
Retiring Director



Estação para o Estudo do Sezonismo, Águas de Moura

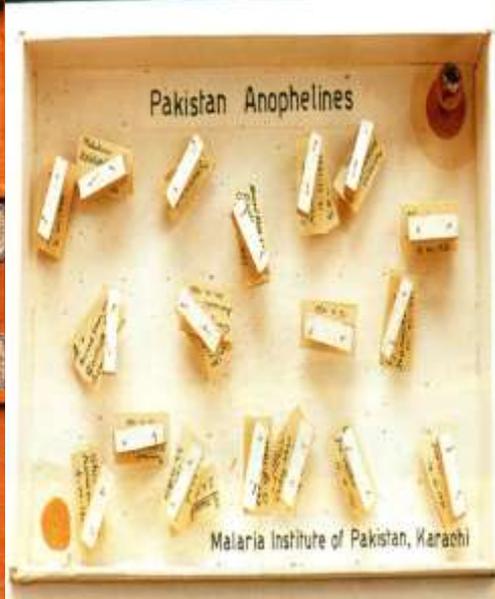
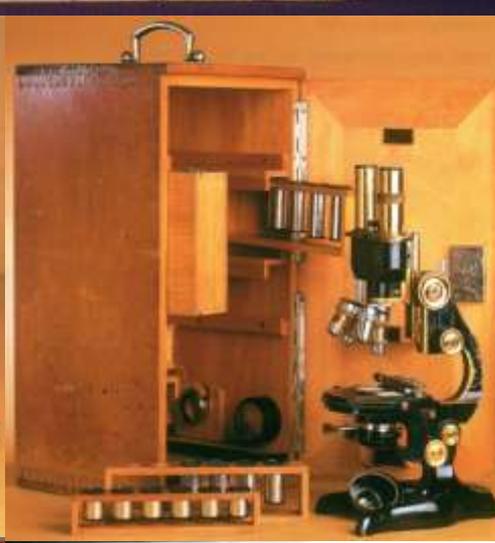
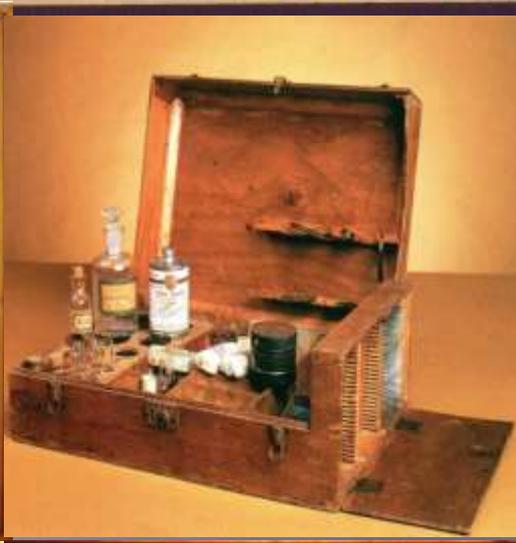


Curso Internacional de Malaria para Médicos. 1952

António Levy Mendes (Portugal), Claude Vermeil (França), Guy Houel (Argélia), J. Fernandez Maruto (Espanha), Branimir Richter (Jugoslávia), Luís Contreras Poza (Espanha), Prof. F. Cambournac, Anastassios Marinou (Grécia), L. Saadat (Irão), George Fameliaris (Grécia), Etienne Noel (França), L. Andarelli (Argélia), Vamona Sinari (Portugal), Jean Deniaud (França), Hector Meyus (Bélgica), Luis Meira (Portugal)



Inauguração do Instituto de Malaria de Águas de Moura  
Fotografia de Américo Ribeiro - Arquivo Municipal de Palmela



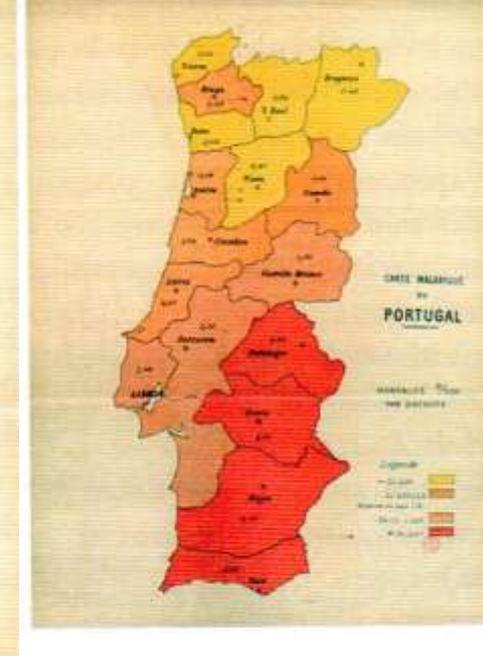
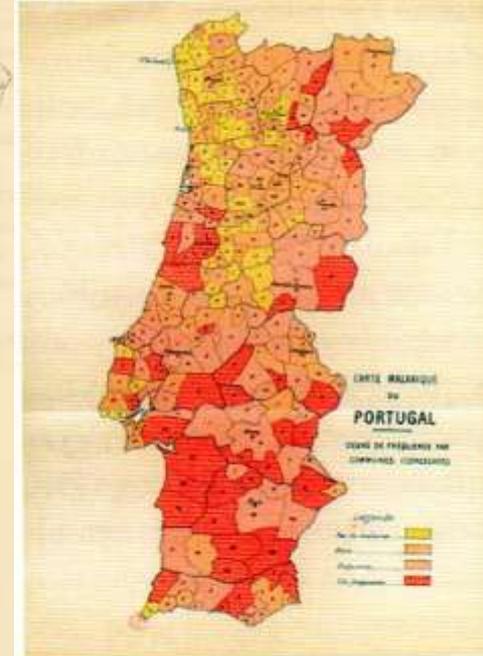
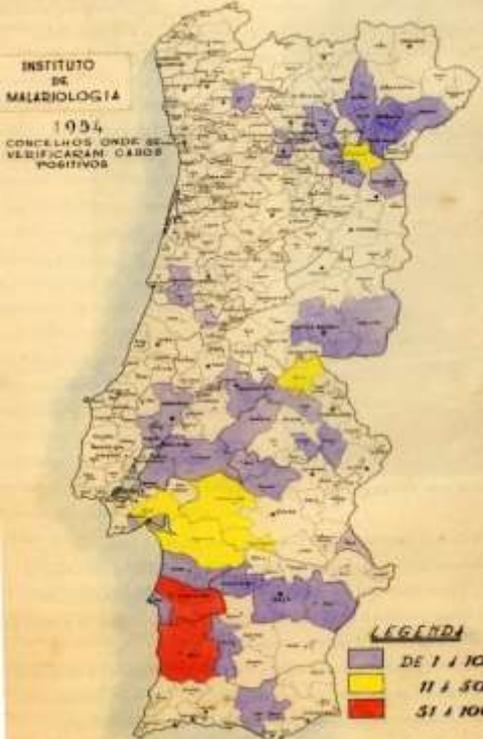


Figura 2 – Carta de sezonismo

Figura 4 – Carta de mortalidade por malária, por distritos

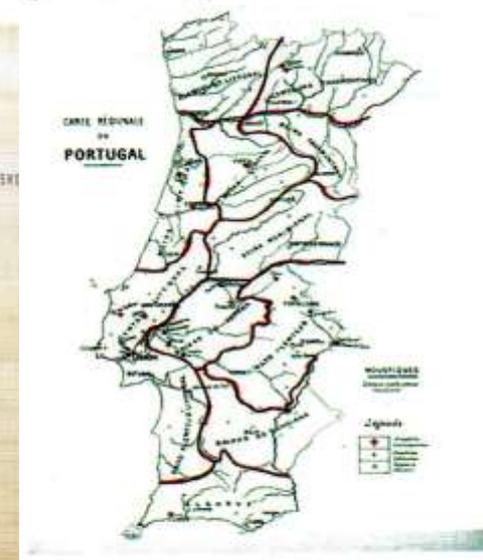
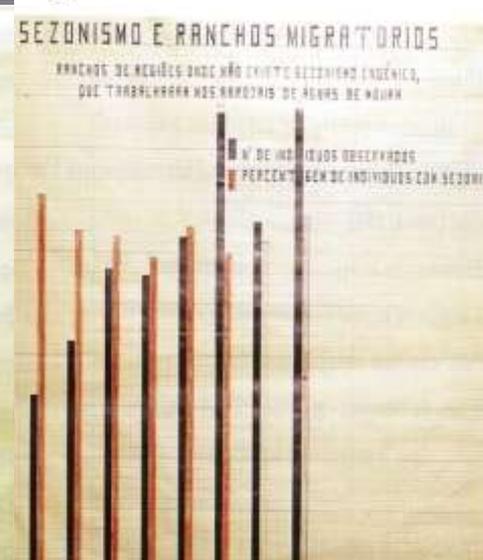
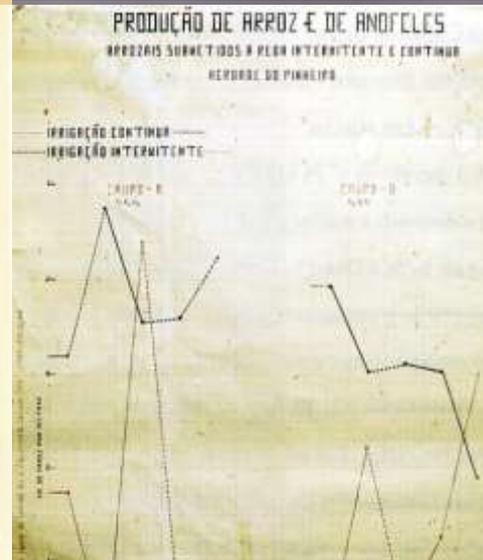
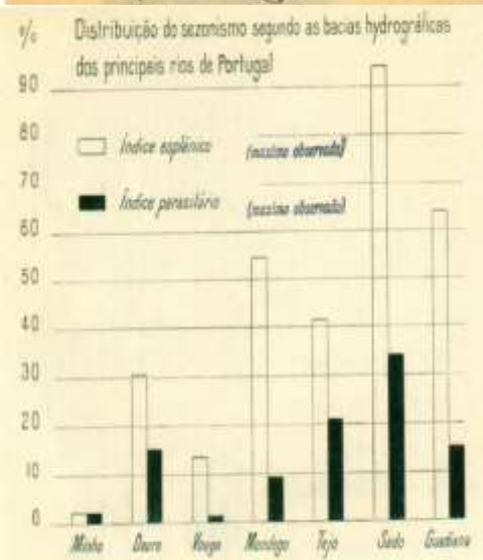


Figura 3 – A distribuição dos vectores da malária em Portugal – Carta de Anophelismo

Figura 3 – A distribuição dos vectores da malária em Portugal – Carta de Anophelismo

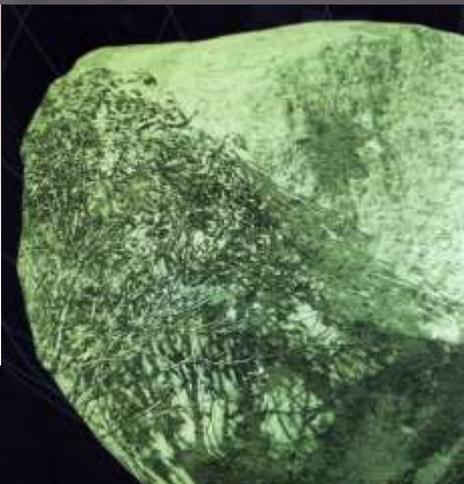
Distribuição do Sezonismo segundo as bacias hidrográficas dos principais rios de Portugal  
in CAMBURNAC, Francisco e LANDERO, Fausto - O Sezonismo em Portugal, 1933



# ESTUDOS NACIONAIS RECENTES NOS DOMÍNIOS DA HISTÓRIA, GEOGRAFIA, ANTROPOLOGIA, EPIDEMIOLOGIA E DA ENTOMOLOGIA

VIAGENS | E  
MISSÕES  
CIENTÍFICAS  
NOS TRÓPICOS

1883 | 2010



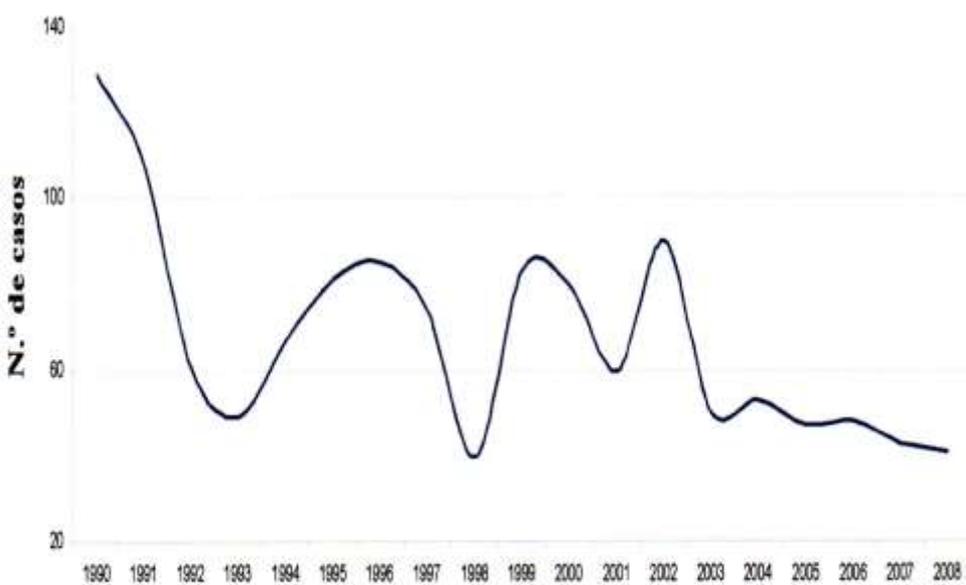


Figura 3.4 – Casos importados de malária em Portugal Continental (1990-2008)  
(fonte dos dados: WHO, 2009)

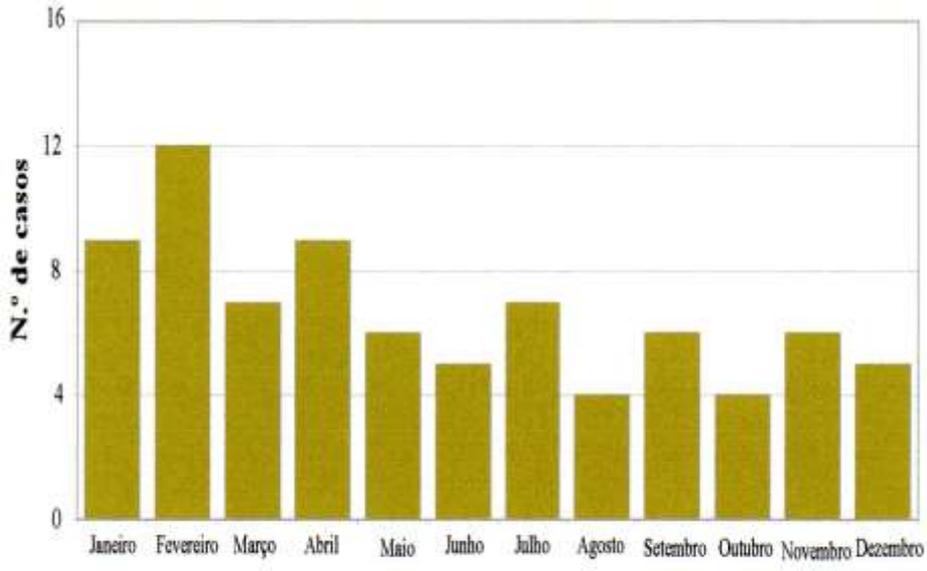


Figura 3.12 – Casos notificados de malária em Portugal Continental, por mês, no ano 2000  
(fonte dos dados: DGS, 2001)

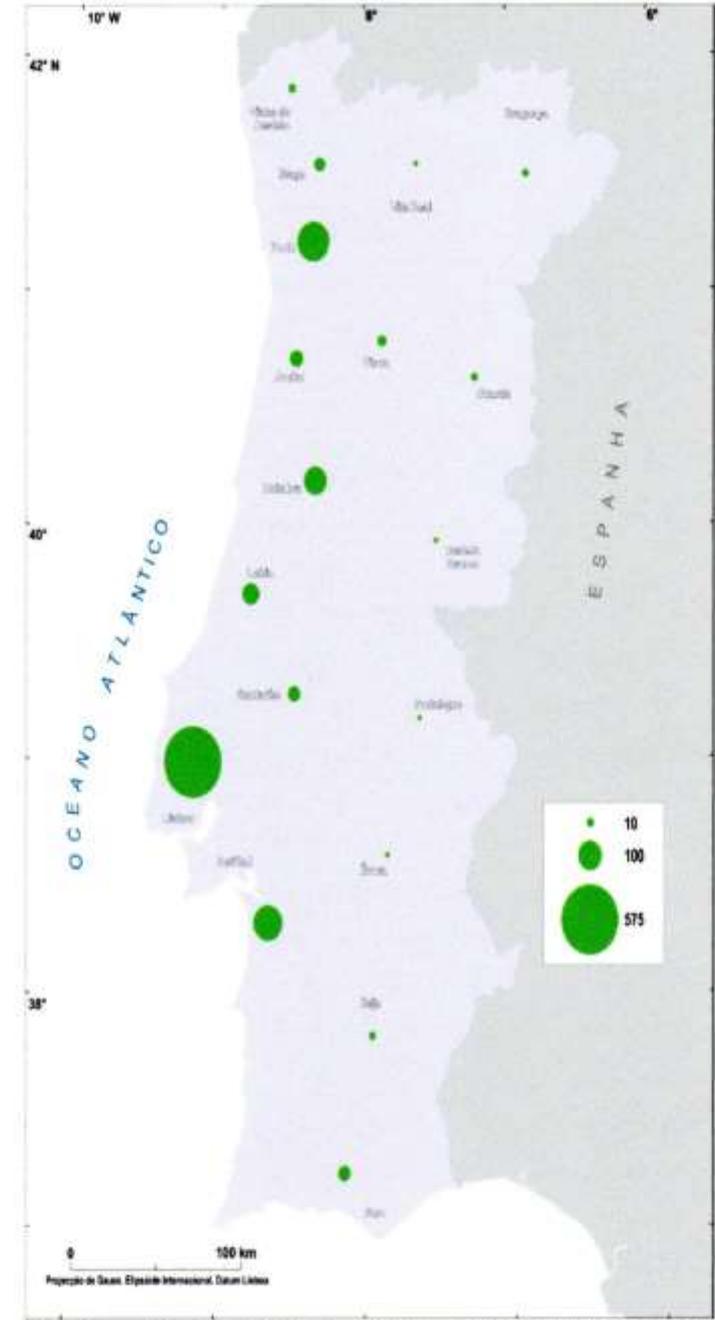


Figura 3.5 – Casos importados de malária, por distrito, em Portugal Continental (total de 1990 a 2006)  
(fonte dos dados: INE, 1991; INE, 1992; INE, 1993; INE, 1994; INE, 1995; INE, 1996; DGS, 2001; DGS, 2002; DGS, 2003; DGS, 2004; DGS, 2007; DGS, 2009)



### “Uma Questão Nacional”

## Enredos da malária em Portugal, séculos XIX e XX



Mónica Alexandra de Almeida Monteiro Saavedra

Doutoramento em Ciências Sociais  
Especialidade: Antropologia Social e Cultural

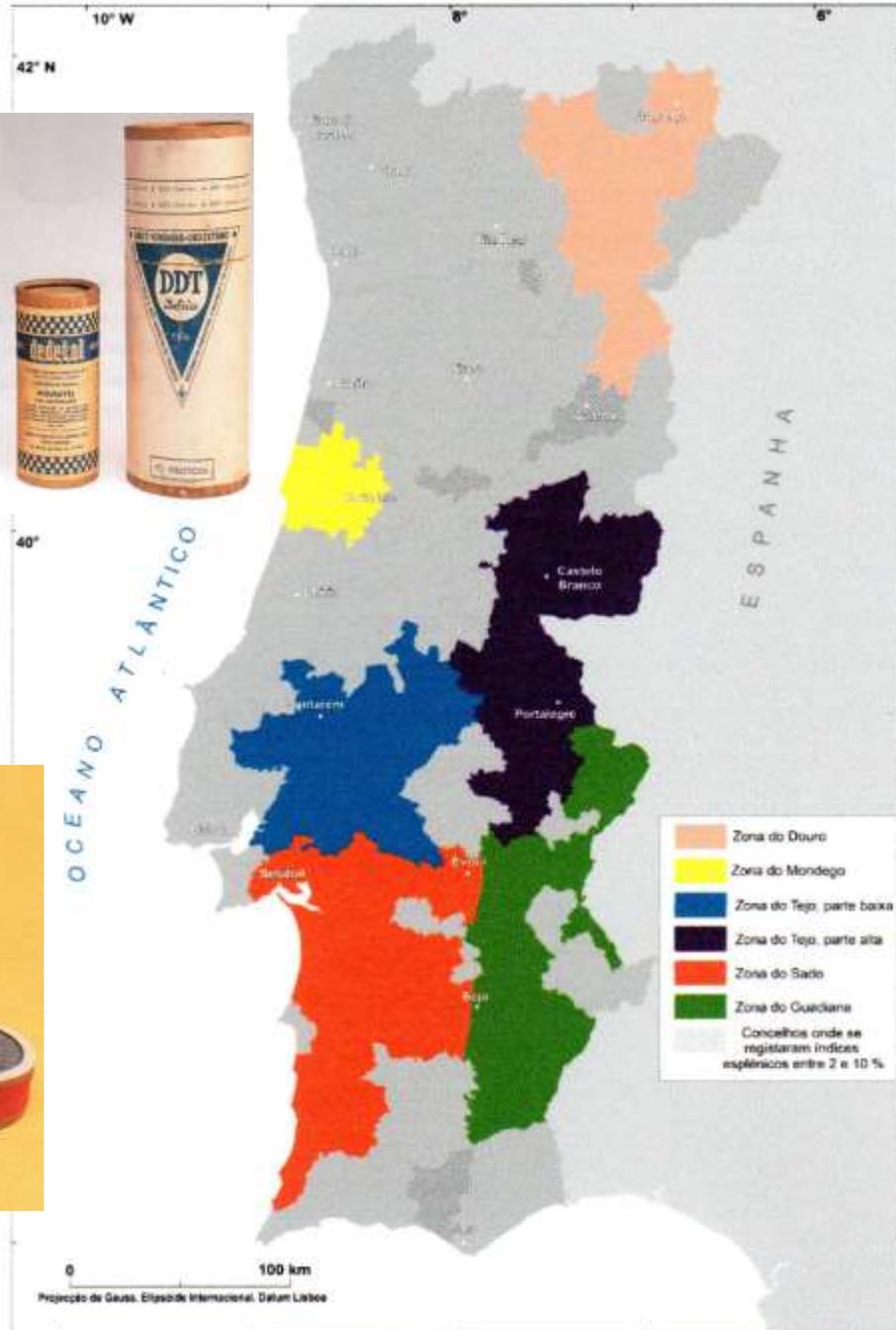


Figura 3.3 – Antigas áreas endémicas de malária em Portugal Continental (adaptado de Cambournac, 1942)

Aquilo, na parte da manhã tava que era uma beleza; tomava o pequeno-almoço, começava a trabalhar, em chegando aí para as onze horas, meio-dia, caía-se um frio comigo, e depois, no fim de resto, um calor... quarenta e não sei quantos graus de febre (Domingos, Azambuja, 2006).

D – Aquela febre, aquela febre que se tinha, o corpo tremia todo.

M – É que a gente queria sustar a tremura e não era capaz; batia mesmo o queixo com a febre (Domingos e Matilde, Azambuja, 2006).

A gente ia para o trabalho [...] ao fim de uma hora ou coisa assim começavam a... em pleno Verão! a tremer todos com um frio terrível e a lançar fora; perdia a força, tínhamos que ir embora para casa (Dâmaso, Vale de Guizo, 2006).

[...] a primeira coisa que aparecia era o frio. Era tremer, não havia roupa que fosse capaz de tapar aquele frio. Depois de passar aquele frio, então é que era largar suor por todo o lado... pronto. E depois, isto era 2-3 dias, ao fim de dois ou três... umas febres de quarenta, quarenta e tal graus sempre que aquilo dava (João, Rio de Moinhos, 2006).

[...] sentia logo aquele frio, aquele frio, aquele frio... e depois passava, depois vinha a febre, começava a ter os lábios tudo arreventado, tudo da febre... bom, no outro dia já a gente sabia, pois passava; aquilo então, passa... tira a vontade de comer mas dura, dura aí uns tantos tempos (João, Rio de Moinhos, 2006).

Aquela sezão era assim: a gente de manhã levantava-se parecia que era capaz de romper Tróia. Quando era daí a bocadinho... aí que malandrice!!! Aquilo não apetecia fazer nada!!! (Francisco, Alcácer do Sal, 2006).

Aquilo durava um dia; no outro dia a gente levantávamos, andávamos... as quartãs era dia sim, dia não; era dia sim, dia não, que era com mais força. E as sezões dava-nos todos os dias; a gente às vezes até nem ia trabalhar (Maria, Vale de Guizo, 2006).

R – [...] havia as sezões, que era dia sim, dia não ou havia as quartãs, que essas eram... as quartãs era quê? de 2 em 2 dias?

D - As quartãs era de 2 em 2 dias mas as sezões... (Rita e Domingas, Benavente, 2005).

Mas é que hoje andávamos boas, a graça era essa, é que hoje andávamos boas e amanhã tavamos com a sezão; e depois era 3 e 4 dias com a sezão; mas quando era os acréscimos era hoje temos amanhã não temos. Chamavam acréscimos: hoje temos, amanhã não temos; no outro dia temos outra vez (Elvira, Benavente, 2005).

M – No outro dia a mema... aquilo era a quase sempre à mema hora

D – Era todos os dias; chegava ali àquela hora, pronto...

M – Chegava ali àquela hora começava a ter febre (Matilde e Domingos, Azambuja, 2006).



Benavente: O seu papel no Combate ao Paludismo em Portugal  
**PALUDISMO BENAVENTE :: 1931-1934**

Dissertação de mestrado orientada por:  
Professor Doutor António Ventura  
Professor Doutor João Cosme

Sandra José Ricardo Ferreira

Direcção Geral de Saúde  
**ESTAÇÃO EXPERIMENTAL**  
DE  
COMBATE AO SEZONISMO  
DE  
**BENAVENTE**

**MANDAMENTOS DA LUTA CONTRA AS SEZDES**

- 1.º - Voto as medidas preventivas que tomo, em casa, quando estou de trabalho, pela a saúde e a do meu lar.
  - 2.º - Não recebo do médico nenhuma prescrição sem a sua ordem.
  - 3.º - Vou sempre a consultas como se fossem com um amigo.
  - 4.º - Frequento a consulta da estação experimental na cidade de Lisboa.
  - 5.º - Quando passo muito tempo no campo que vivo e as larvas das sezdes que encontro.
  - 6.º - Nunca desisto de ao primeiro tratamento não se curar, paralizo a todo o caso.
  - 7.º - Não deixo ficar sem obreção a outros e as minhas.
  - 8.º - Não deixo de convencer a meu vizinho a vizinha e a família.
  - 9.º - Não deixo dormir com as janelas abertas se elas não tiverem insecticidas.
  - 10.º - Não cultivo a produtividade de plantas péras no campo com vegetação desenvolvida.
- Estas são as medidas tomadas em casa. Tomar regularmente a quina e seguir fielmente as ordens do médico.

Doc. 1  
Propaganda de Propaganda, 1931 (Plano para Estação Experimental de Benavente)



Mapa 1  
Localização geográfica do município de Benavente com indicação das respectivas freguesias (2008)

# A APLICAÇÃO DE SISTEMAS DE INFORMAÇÃO GEOGRÁFICA NA MODELAÇÃO DA DISTRIBUIÇÃO DE VECTORES DE DOENÇAS: O MOSQUITO VECTOR DA MALÁRIA *ANOPHELES ATROPARVUS* EM PORTUGAL CONTINENTAL

UNIVERSIDADE DE LISBOA  
FACULDADE DE LETRAS  
DEPARTAMENTO DE GEOGRAFIA

César Capinha<sup>1</sup>; Eduardo Gomes<sup>2</sup>; Eusébio Reis<sup>1</sup>; Jorge Rocha<sup>1</sup>; Carla A. Sousa<sup>2</sup>; Virgílio E. do Rosário<sup>3</sup>; A. Paulo G. Almeida<sup>2</sup>

<sup>1</sup> Centro de Estudos Geográficos – Universidade de Lisboa, Alameda da Universidade, 1600-214 Lisboa

<sup>2</sup> Unidade de Entomologia Médica – Instituto de Higiene e Medicina Tropical – Universidade Nova de Lisboa, Rua da Junqueira, 96, 1349-008 Lisboa

<sup>3</sup> CMDT-LA – Instituto de Higiene e Medicina Tropical – Universidade Nova de Lisboa, Rua da Junqueira, 96, 1349-008 Lisboa

(cesarcapinha@hotmail.com; eduardojonas@gmail.com; jrocha@fl.ul.pt; eusebioreis@campus.ul.pt; casousa@ihmt.unl.pt; virgilio.rosario@ihmt.unl.pt; palmeida@ihmt.unl.pt)



## RESUMO

A malária constituiu um dos principais problemas de saúde pública da primeira metade do séc. XX em Portugal. Actualmente, embora não se registem casos autóctones, a ocorrência constante de casos importados, associados a uma potencial capacidade de transmissão dos seus agentes infecciosos (plasmódios) pelo mosquito *Anopheles atroparvus*, mantém presente o risco de futuras infecções endémicas. No sentido de melhor se conhecer a distribuição desta espécie em Portugal Continental foram efectuados diversos modelos de distribuição potencial desta espécie vectora. Com base em vários levantamentos de campo referentes à presença de espécies do complexo *Anopheles maculipennis* e diversos factores ambientais considerados influentes na sua distribuição, foram utilizados cinco métodos correlativos distintos: distância de Mahalanobis; rede neuronal artificial; regressão logística binária; máxima entropia e algoritmo genético. A validação efectuada através do índice *kappa* permitiu identificar os modelos provenientes da regressão logística binária e máxima entropia como os de melhor desempenho preditivo. Com vista à redução da incerteza, os cinco modelos foram posteriormente combinados sob a forma de mapas binários. O seu resultado foi confrontado com as principais áreas de incidência de malária em Portugal na primeira metade do séc. XX, tendo-se verificado uma elevada concordância espacial entre as duas representações.

O MOSQUITO VECTOR DA MALÁRIA *ANOPHELES ATROPARVUS*, VAN THIEL, 1927: ADEQUABILIDADE DE *HABITAT* EM PORTUGAL CONTINENTAL E POTENCIAIS ALTERAÇÕES FUTURAS DO SEU ESPAÇO CLIMÁTICO

César Dinis Santos Capinha

Mestrado em Sistemas de Informação Geográfica e Modelação Territorial Aplicados ao Ordenamento

# A Importância determinante Factores Meteriologicos

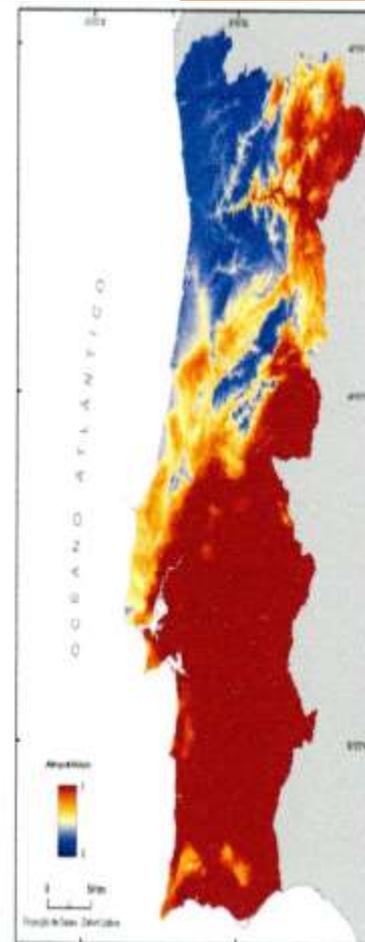
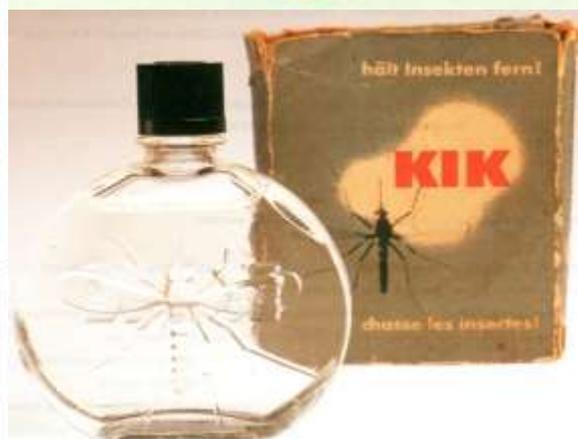
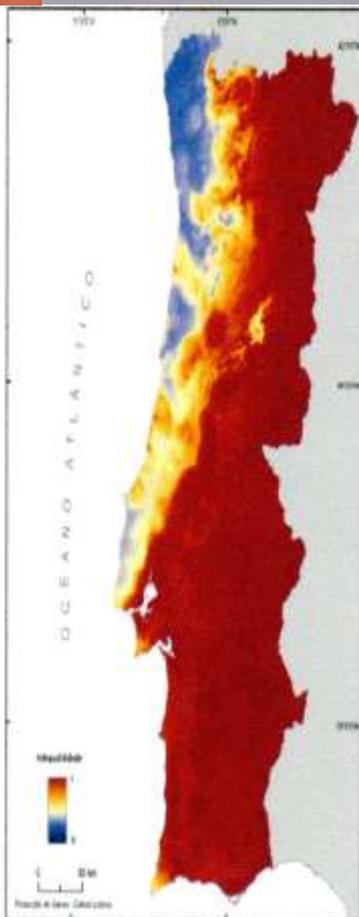


Figura 5 – Adequabilidade de *habitat* para *Anopheles atroparvus* com base na regressão logística binária Figura 6 – Adequabilidade de *habitat* para *Anopheles atroparvus* com base em redes neuronais artificiais

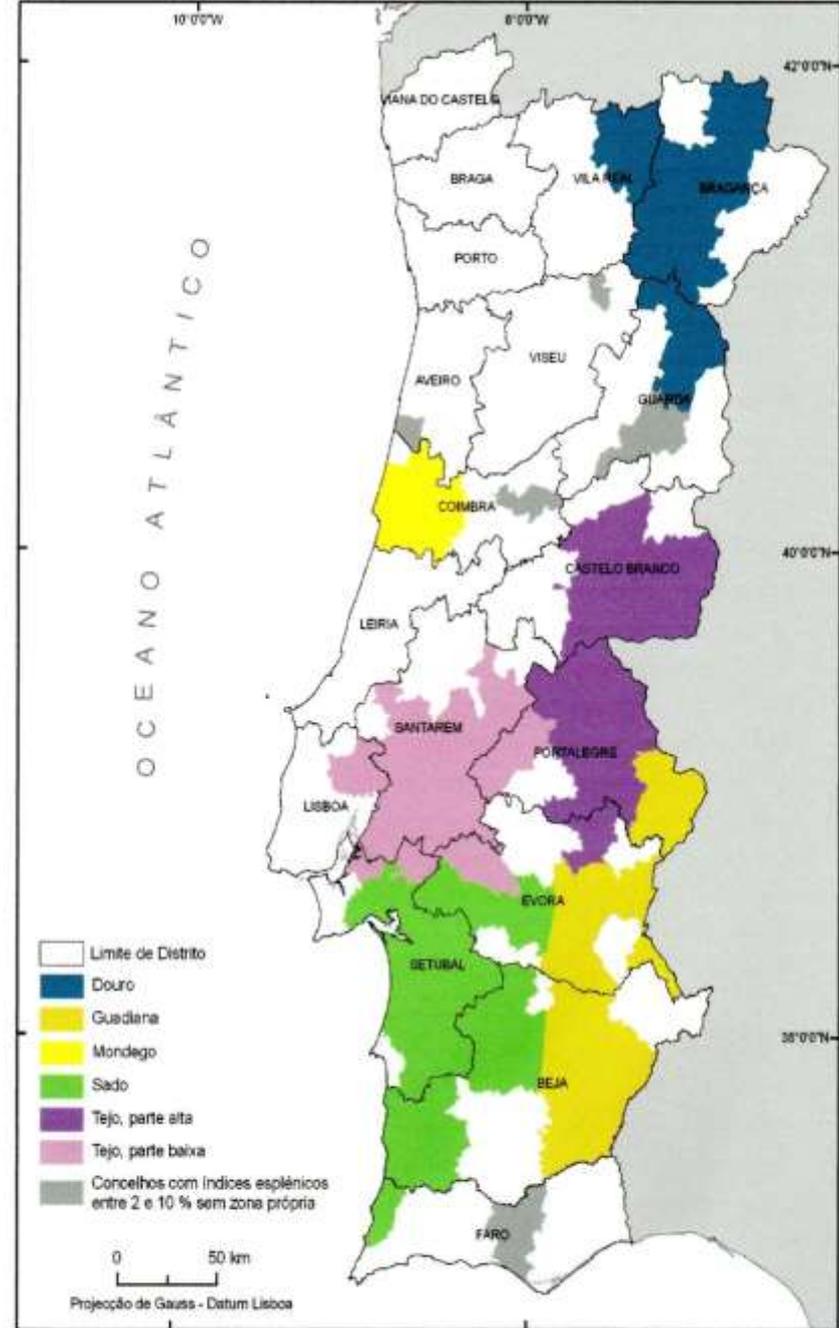
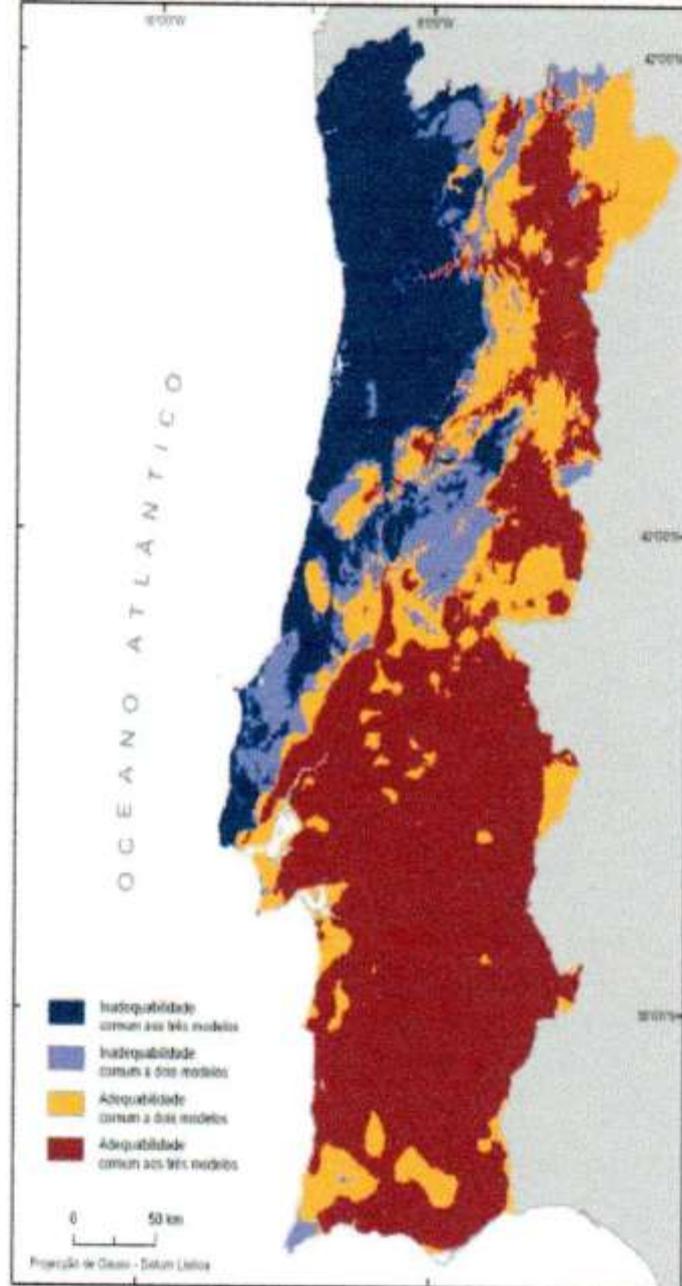


Figura 7 – Combinação de modelos binários de adequabilidade

Figura 6a - Zonas de malária em Portugal na primeira metade do séc. XX (adaptado de Cambournac, 1942)

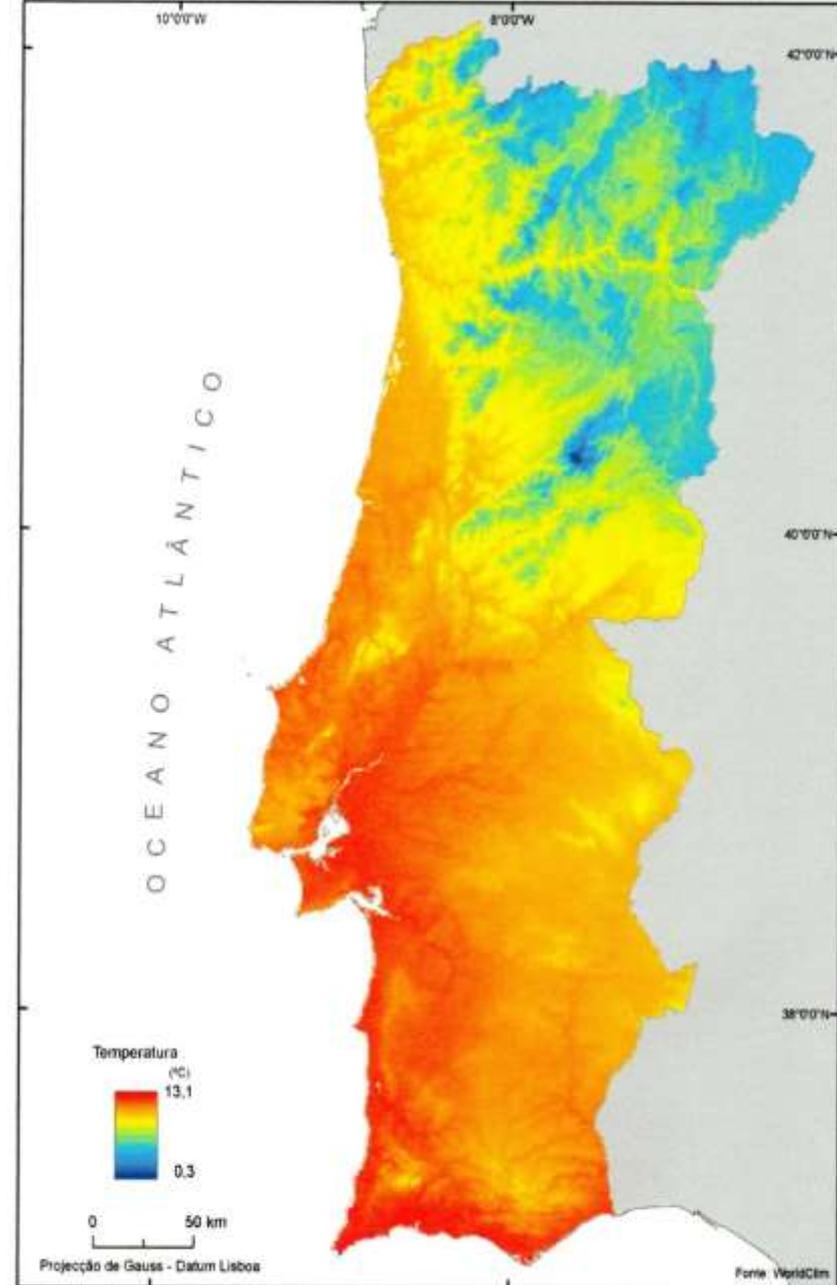
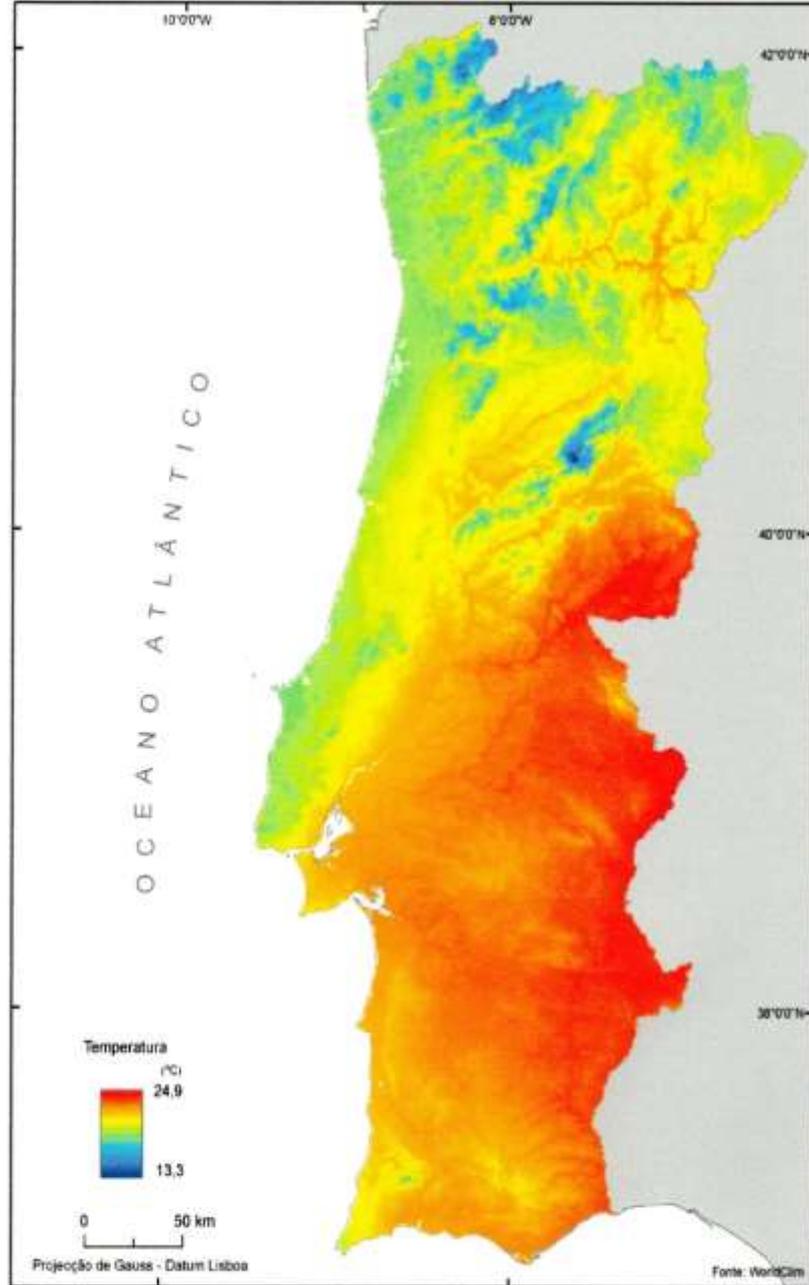


Figura 1a – Temperatura média do trimestre mais quente em Portugal Continental para o período 1950-2000    Figura 2a – Temperatura média do trimestre mais frio em Portugal Continental para o período 1950-2000

(Fonte: Projecto WorldClim)

(Fonte: Projecto WorldClim)

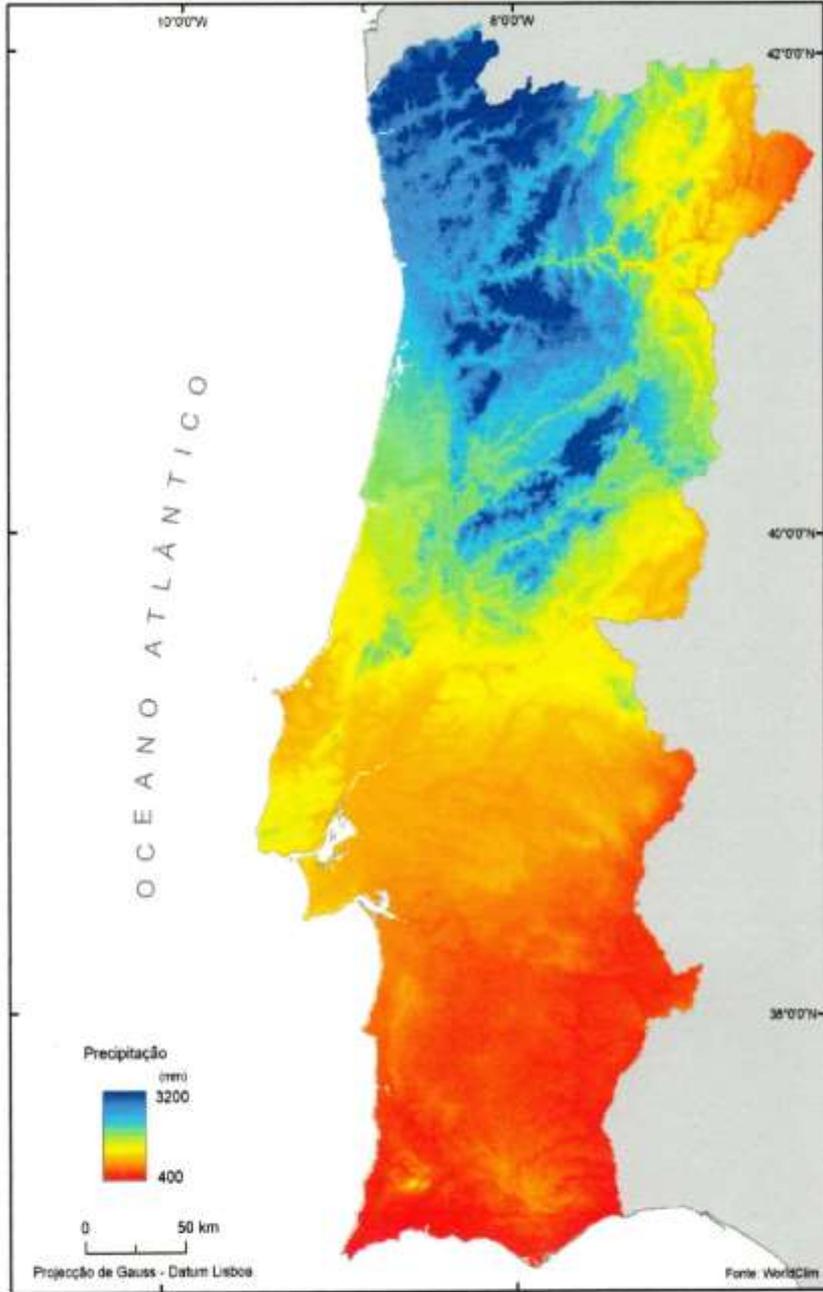


Figura 3a – Precipitação anual média em Portugal Continental para o período 1950-2000 (Fonte: Projecto WorldClim)

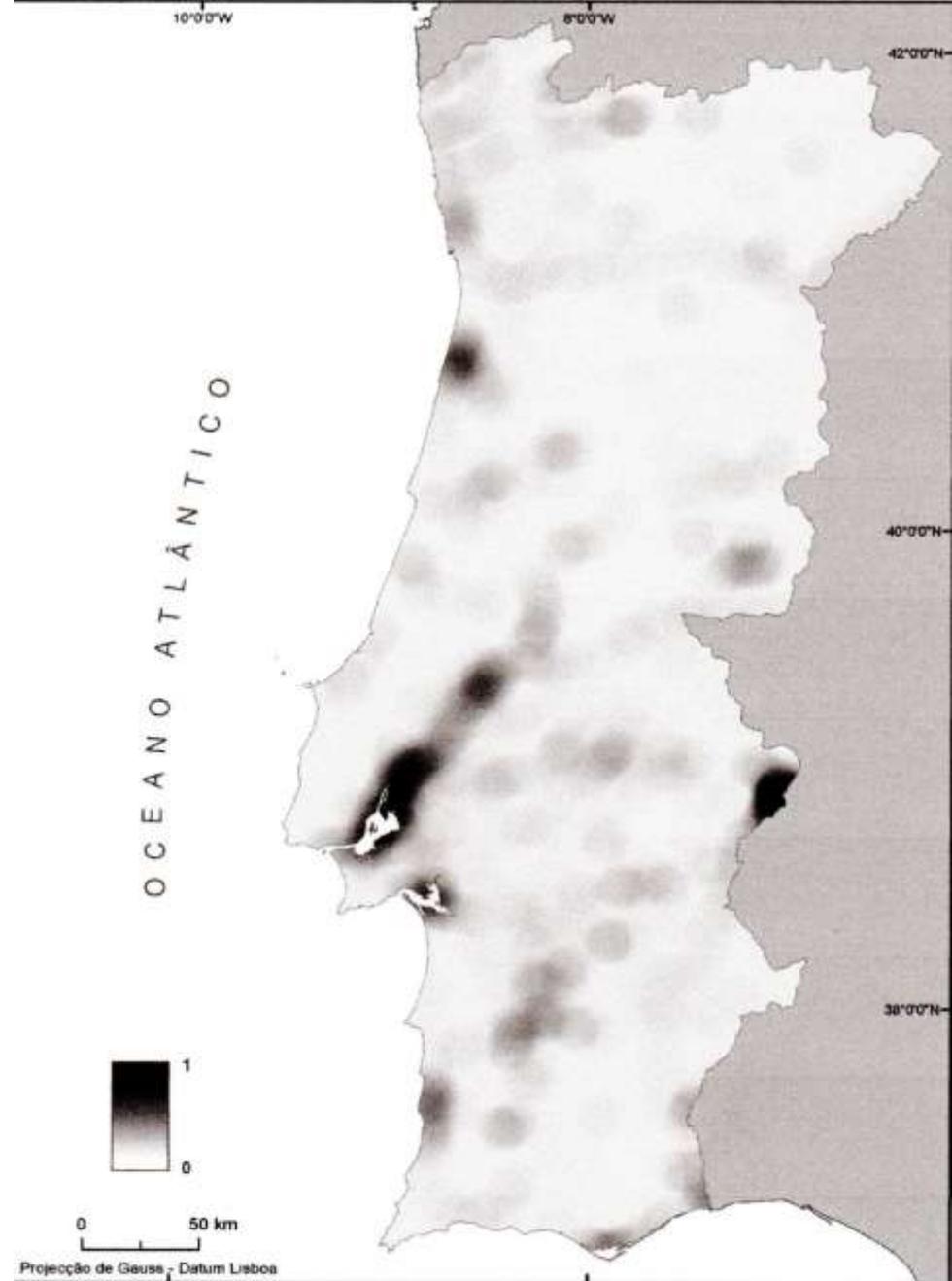


Figura 4a – Índice de disponibilidade e adequabilidade de zonas húmidas

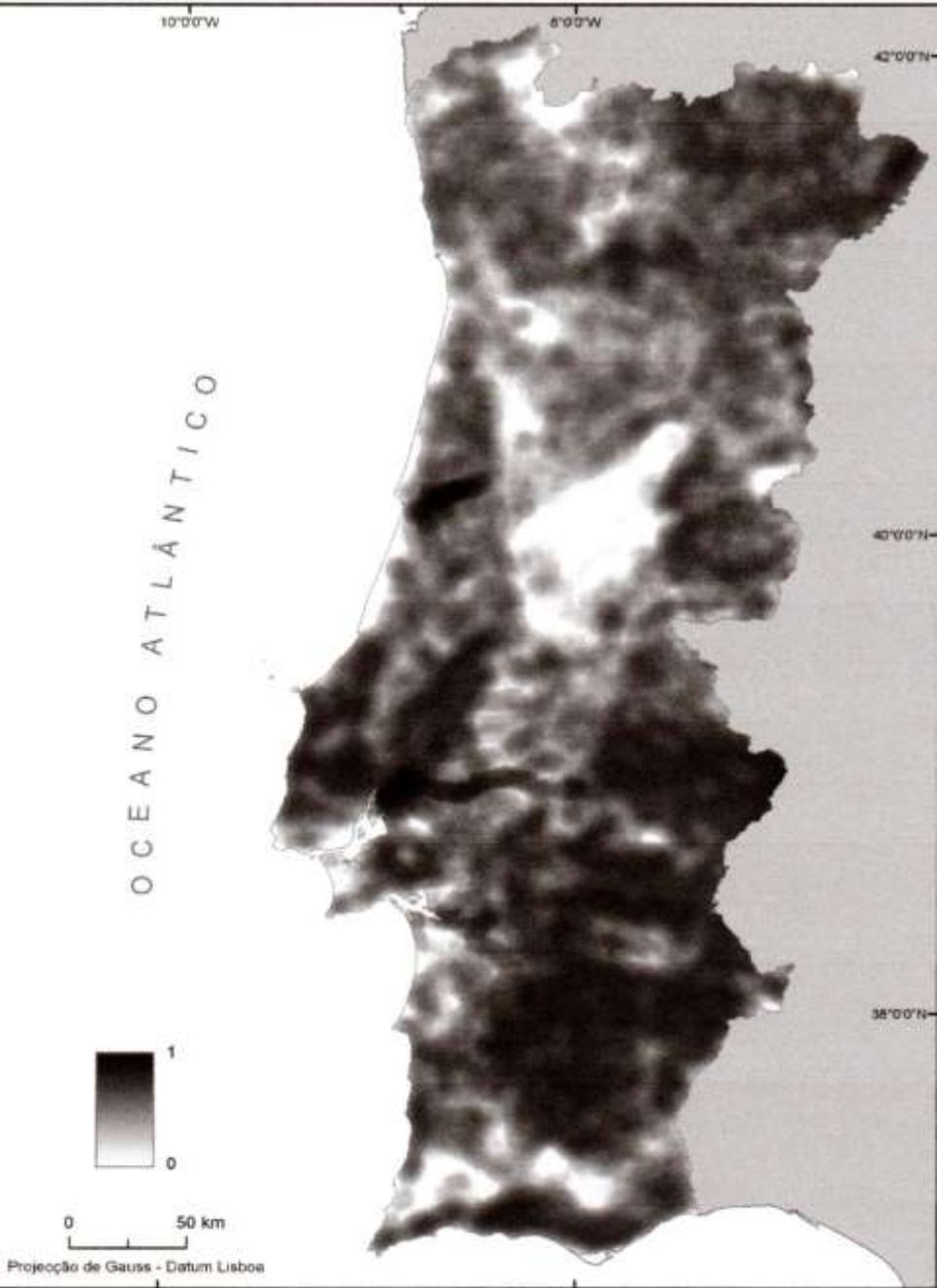


Figura 5a – Índice de disponibilidade e adequabilidade de uso de solo agrícola

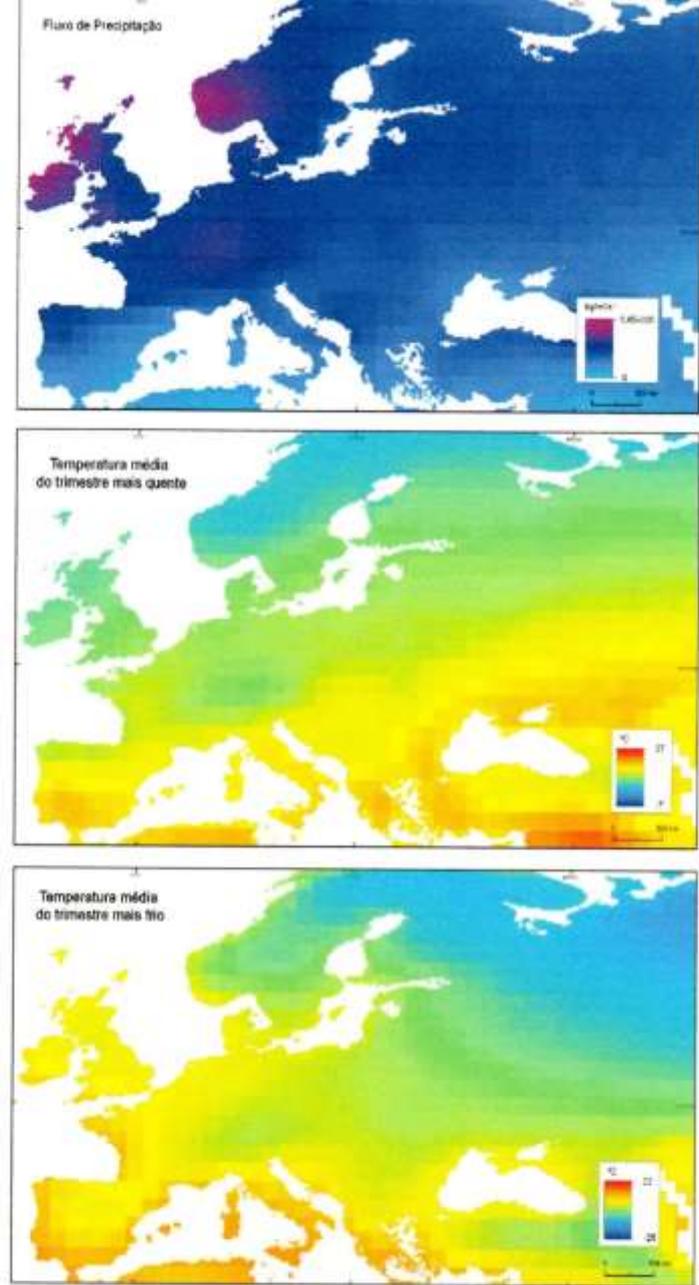


Figura 7a – Precipitação e temperatura da média de experimentações 20th Century Experiment do modelo CCSM-3 para o período 1970-2000

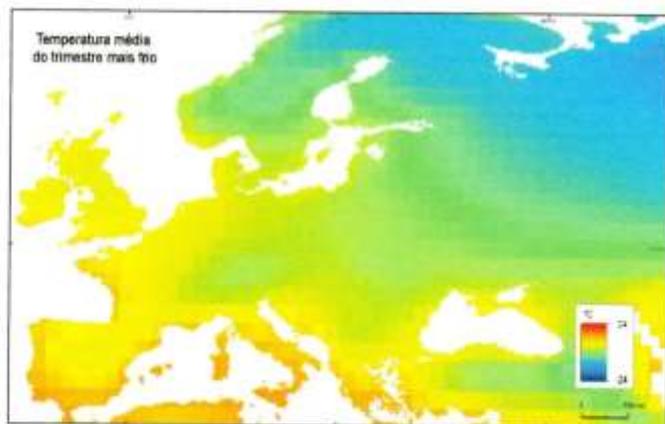
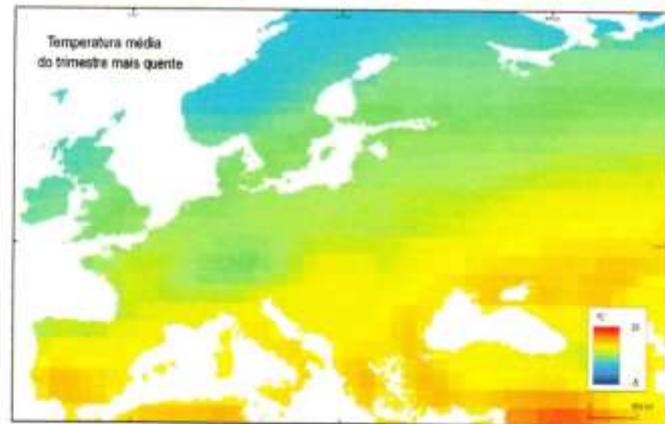
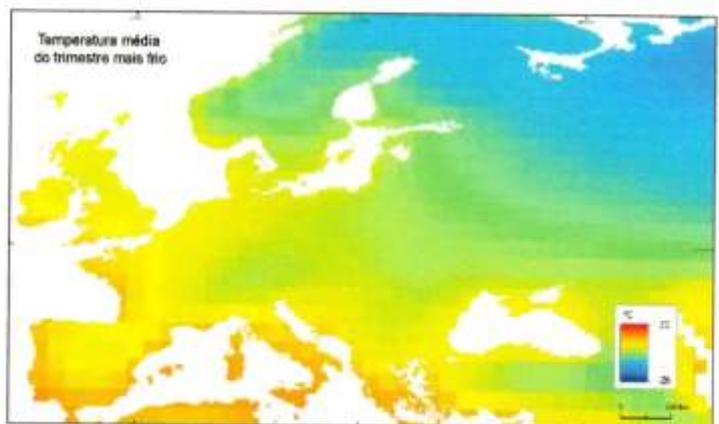
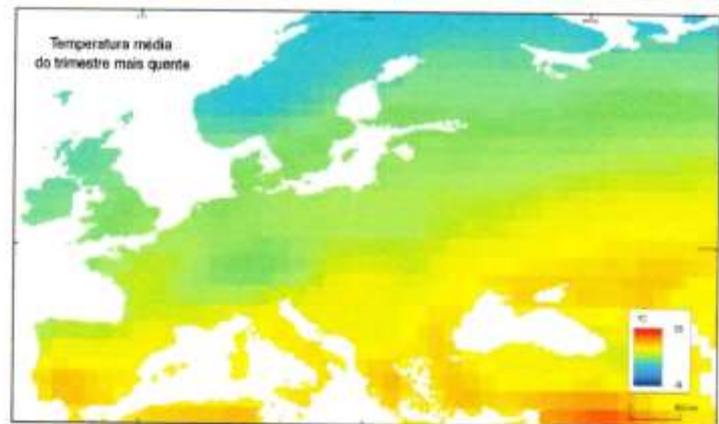
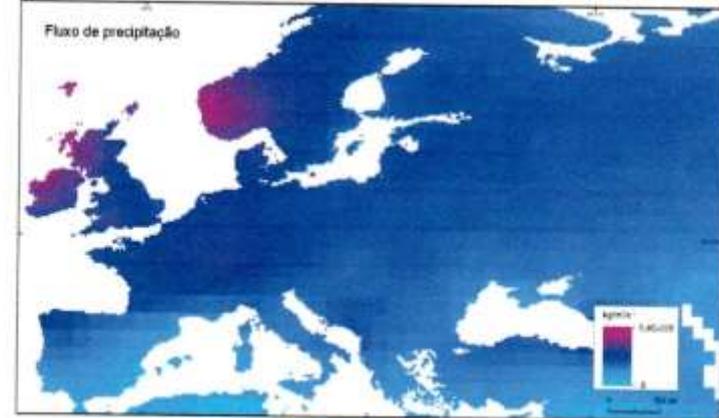


Figura 8a – Precipitação e temperatura da média de experimentações *Climate Change Commitment* do modelo CCSM-3 para o período 2040-2070

Figura 9a – Precipitação e temperatura da média de experimentações do cenário B1 do modelo CCSM-3 para o período 2040-2070

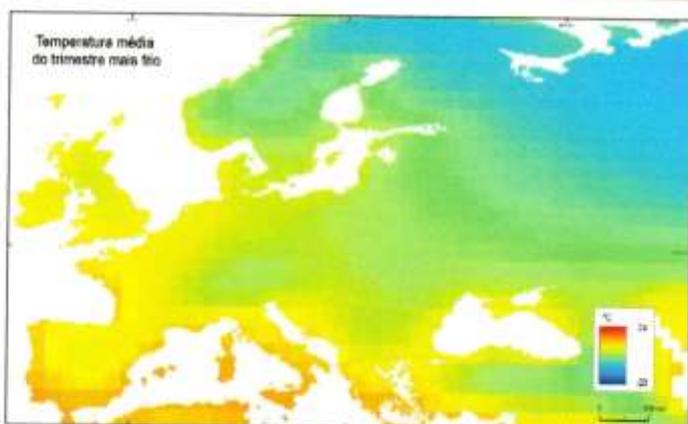
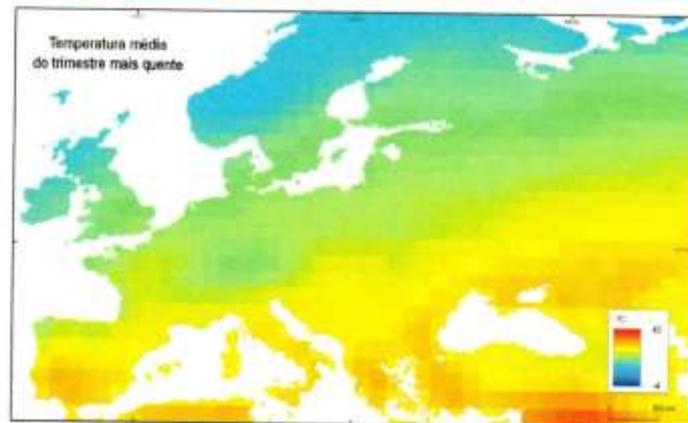
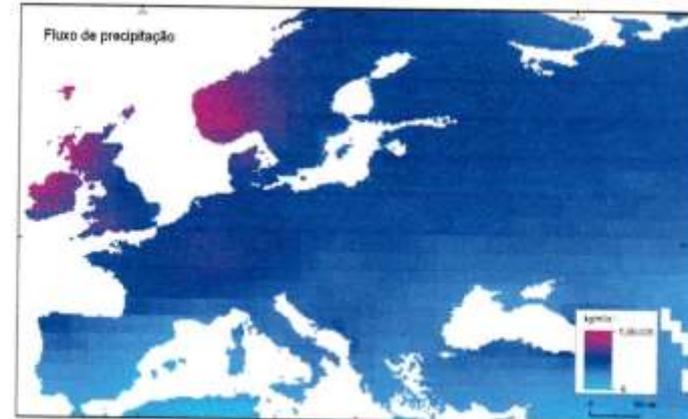
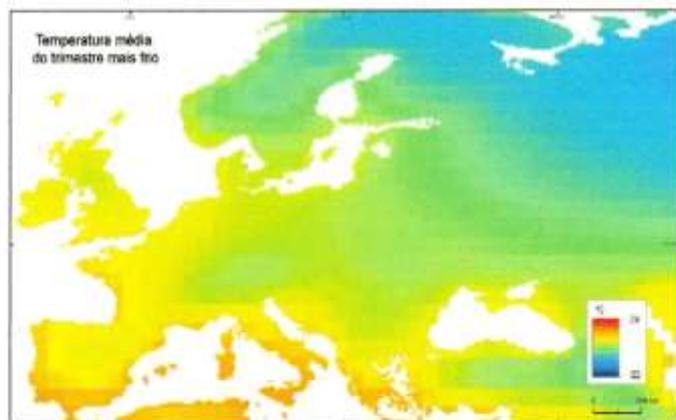
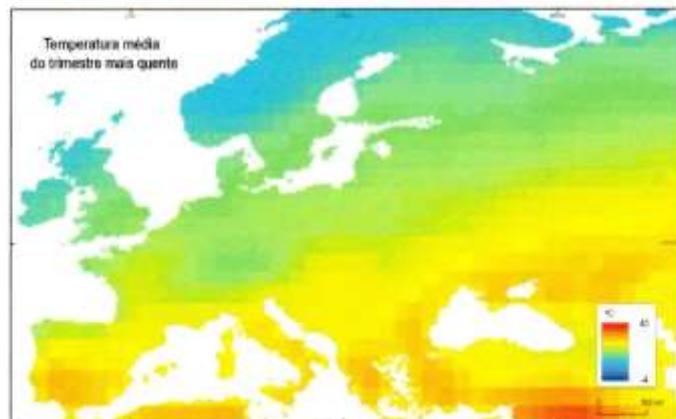


Figura 10a – Precipitação e temperatura da média de experimentações do cenário A1B do modelo CCSM-3 para o período 2040-2070

Figura 11a – Precipitação e temperatura da média de experimentações do cenário A2 do modelo CCSM-3 para o período 2040-2070

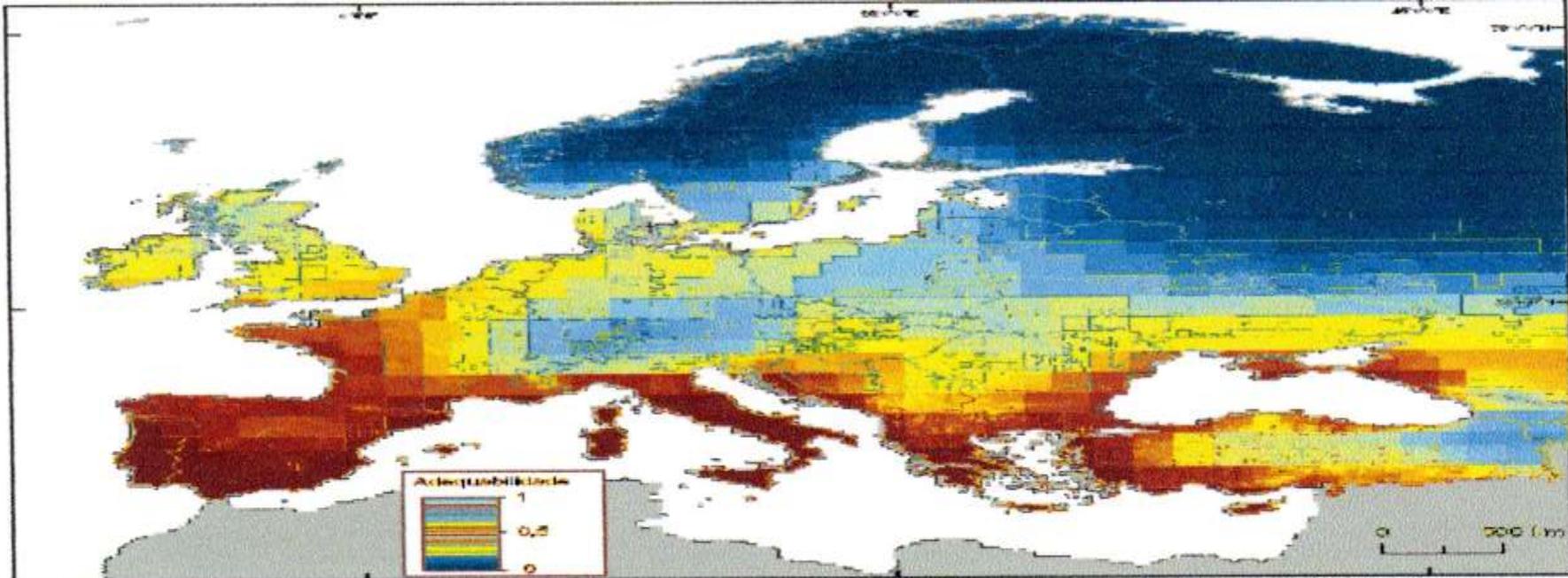


Figura 12 – Modelo de espaço climático actual para *Anopheles atroparvus*

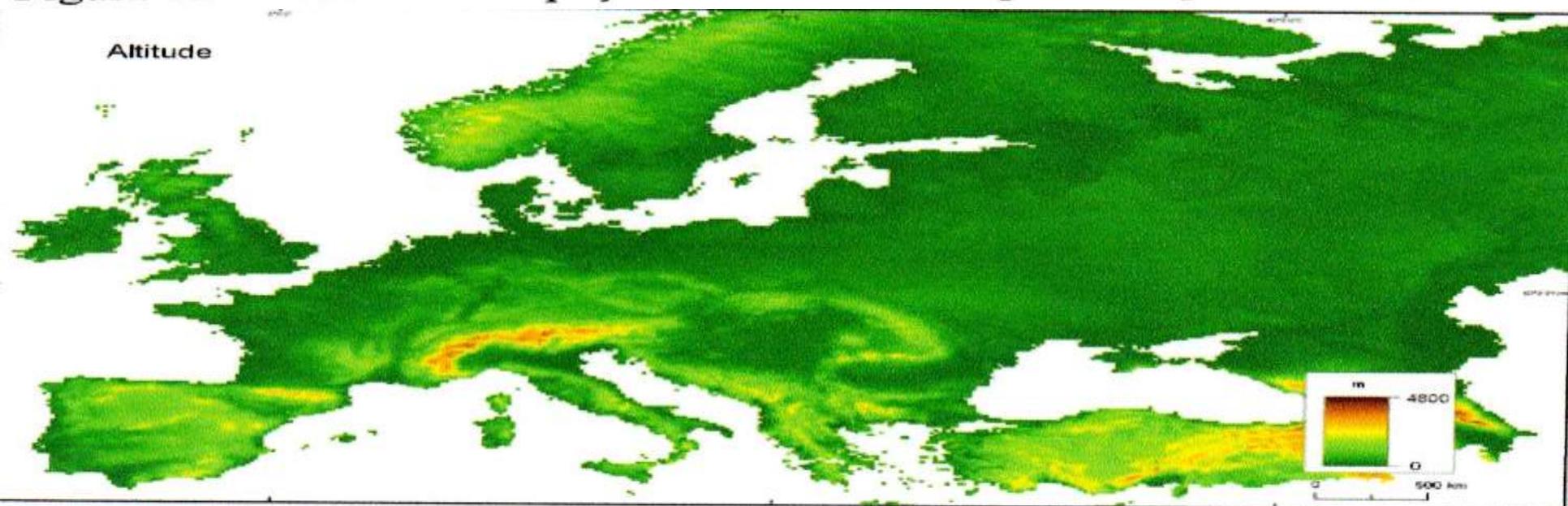


Figura 12a – Modelo de altitudes para o continente Europeu

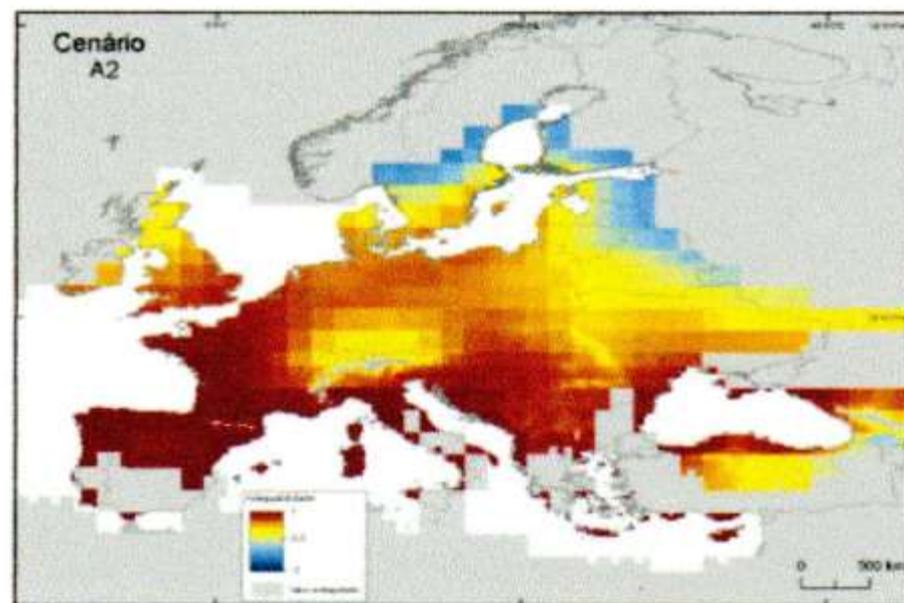
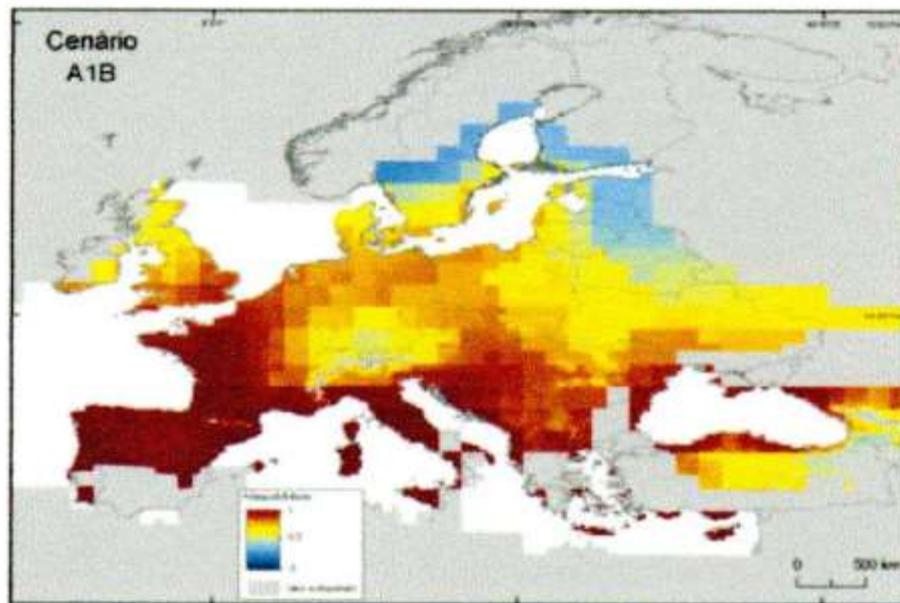
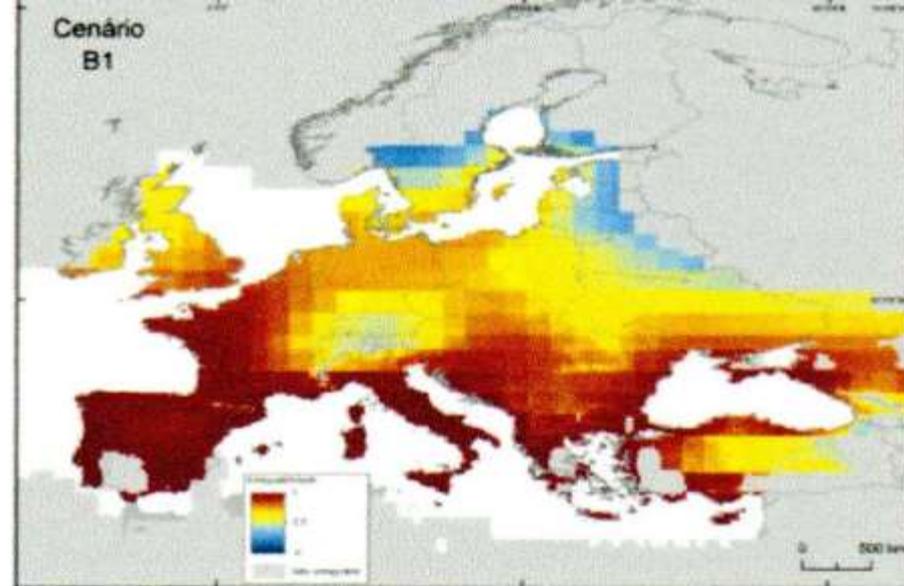
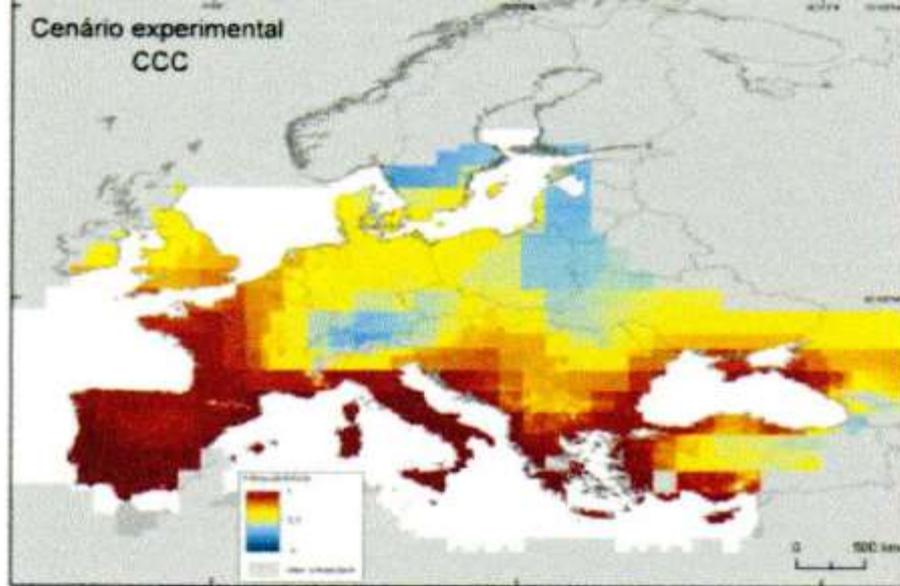


Figura 13 – Espaços climáticos de *Anopheles atroparvus* com base em cenários climáticos futuros para o período 2040 - 2070

# RISCO POTENCIAL DE TRANSMISSÃO DE MALÁRIA EM PORTUGAL CONTINENTAL

Eduardo Jonas da Costa Gomes

Dissertação de Mestrado em Gestão do Território – Área de Especialização em Detecção Remota e Sistemas de Informação Geográfica

JANEIRO, 2010



Criança de 3 anos, Águas de Moura  
Infeção de *Pl. vivax* - Baço 4 da escala de Boyd



Cabanas de trabalhadores, Zambújal, c. 1935

Figura 2.1 – Presenças e ausências de *Anopheles atroparvus* em Portugal Continental

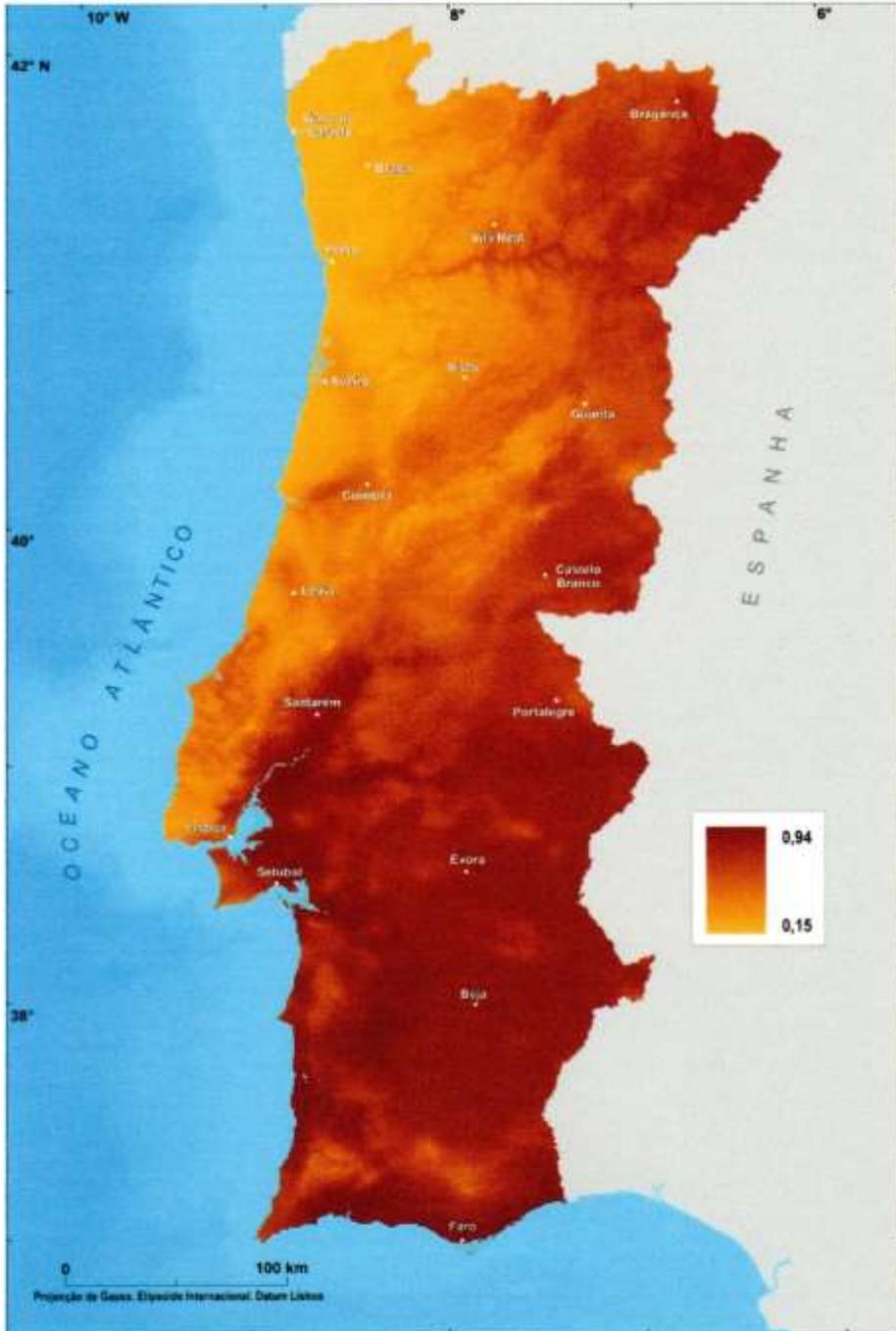


Figura 2.5 – Modelo preditivo de adequabilidade de habitat de *Anopheles atroparvus* para Portugal Continental

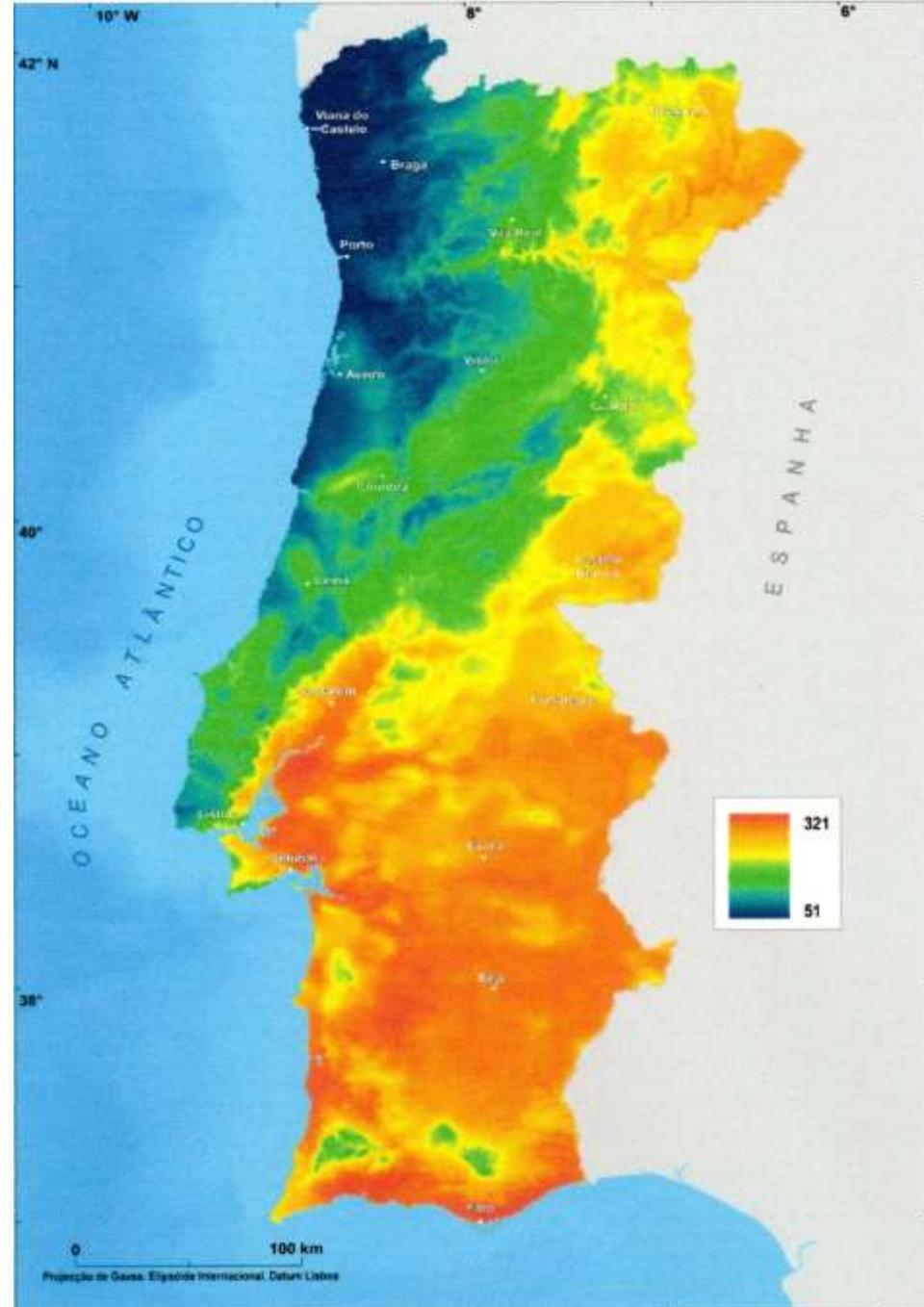


Figura 2.7 – Modelo preditivo da abundância de *Anopheles atroparvus* para Portugal Continental

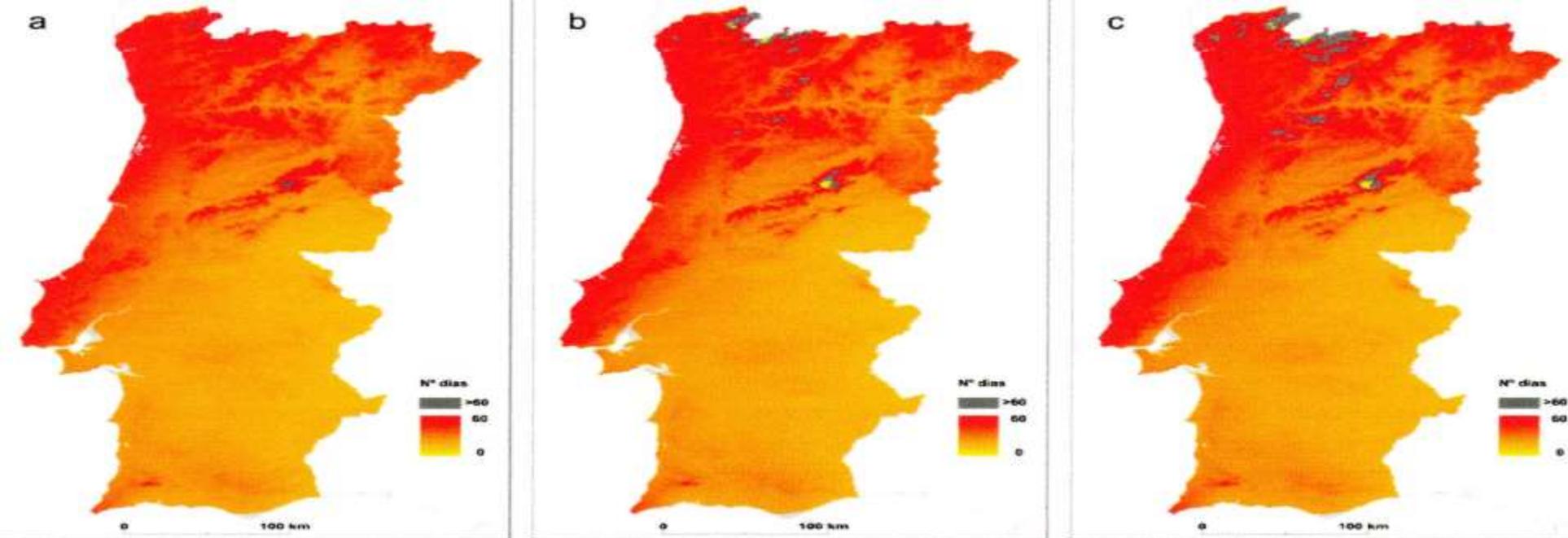


Figura 3.9 – Número de dias em que os 3 tipos de plasmódios se desenvolvem no mês de Agosto: *Plasmodium vivax* (a); *Plasmodium falciparum* (b) e *Plasmodium malariae* (c)

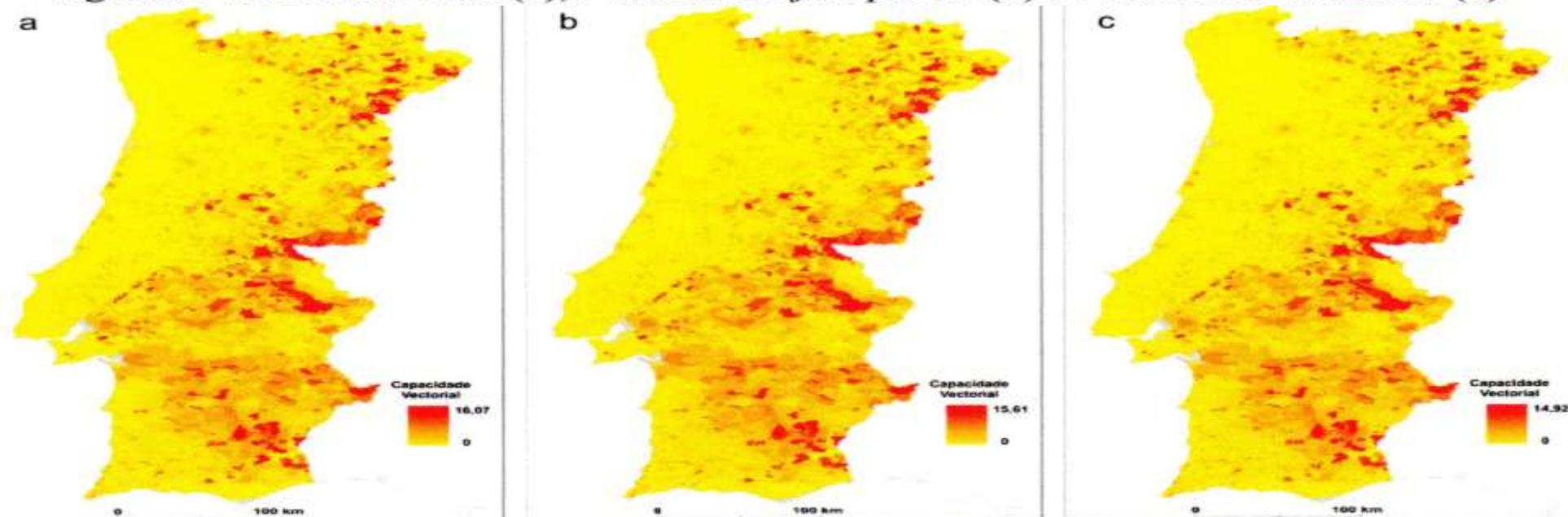


Figura 3.10 – Índice de capacidade vectorial para *Anopheles atroparvus*, com receptividade a *Plasmodium vivax* (a); *Plasmodium falciparum* (b) e *Plasmodium malariae* (c)

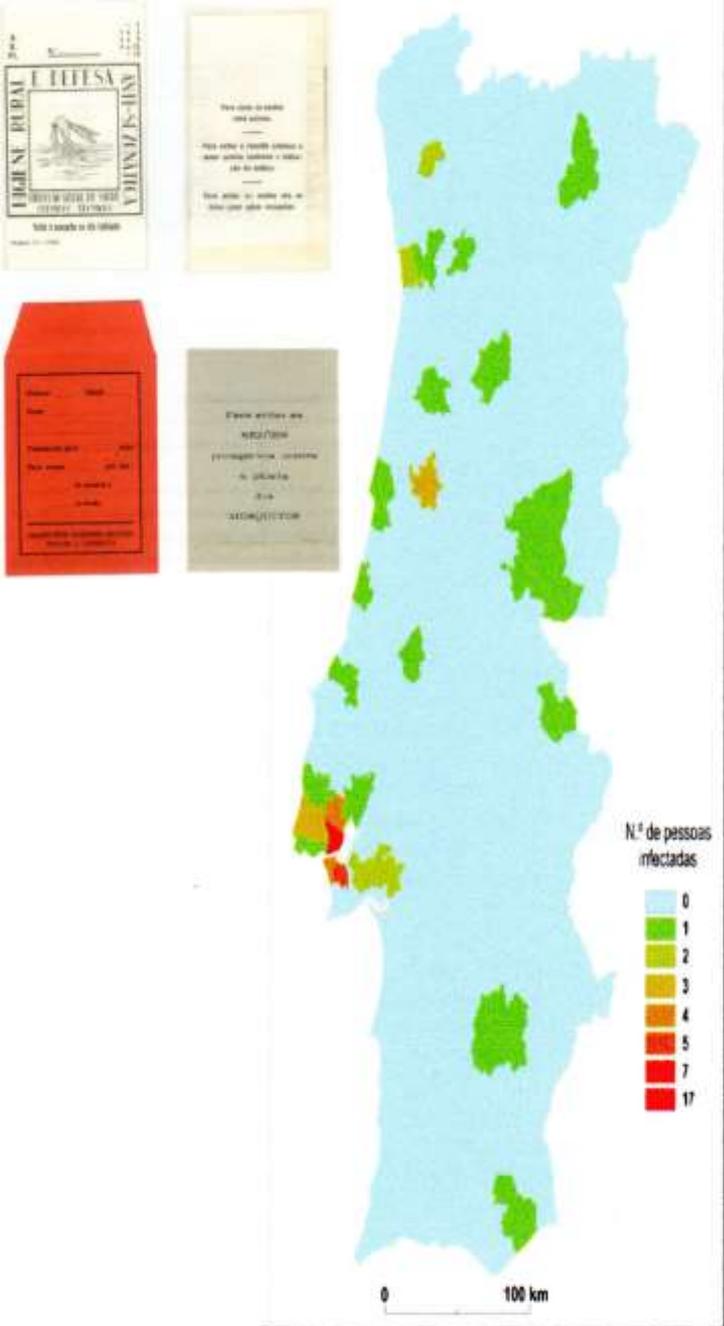
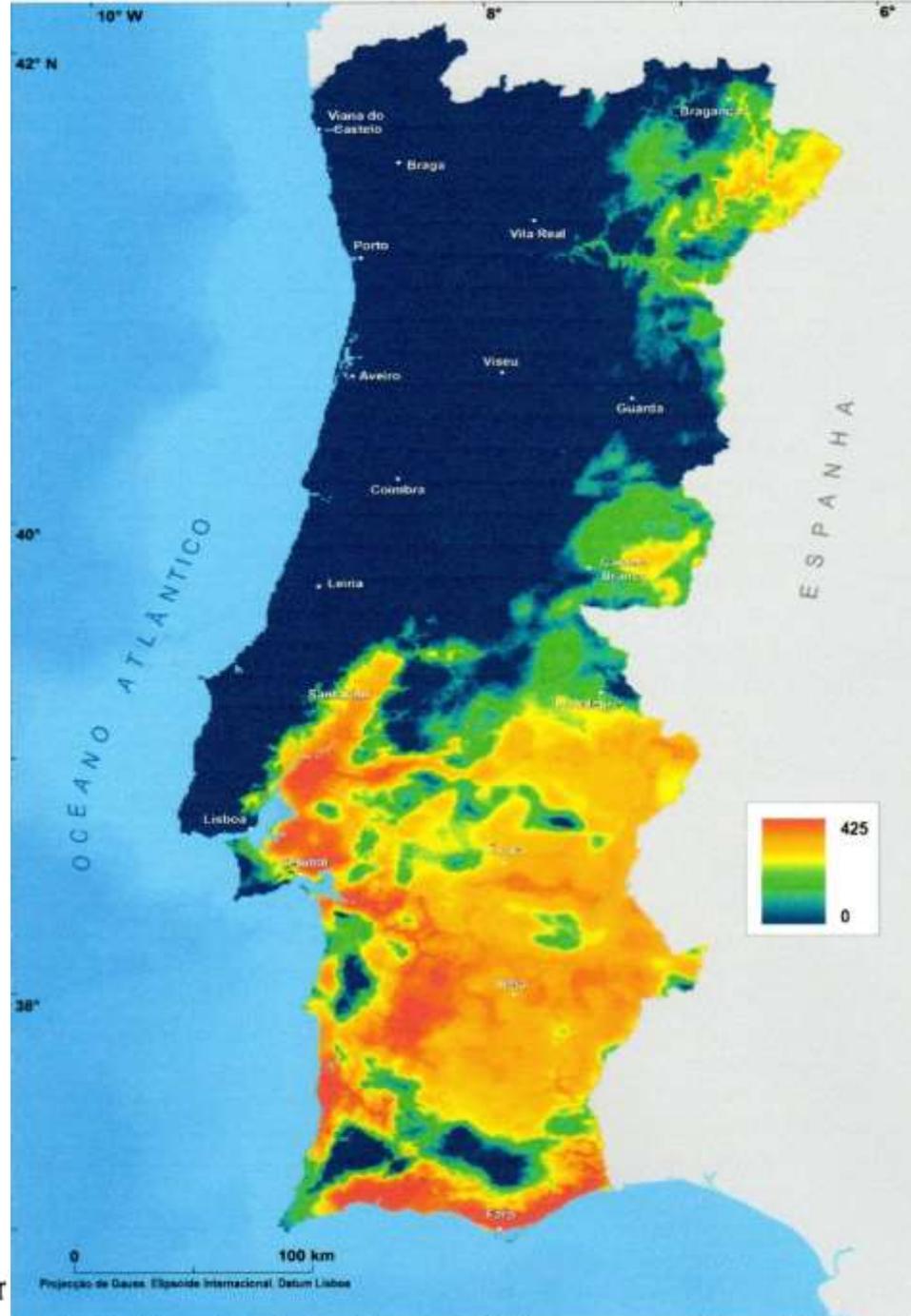
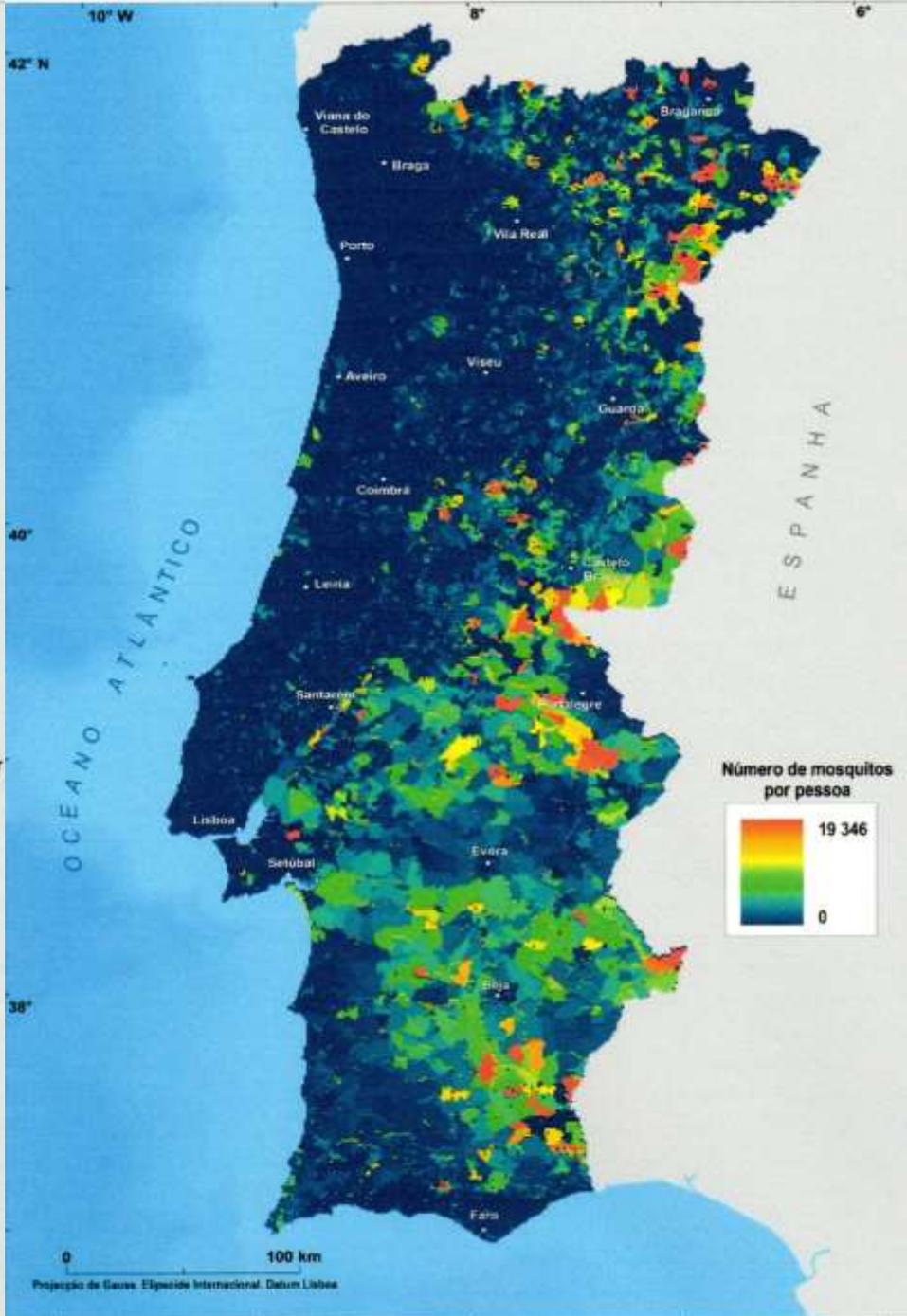
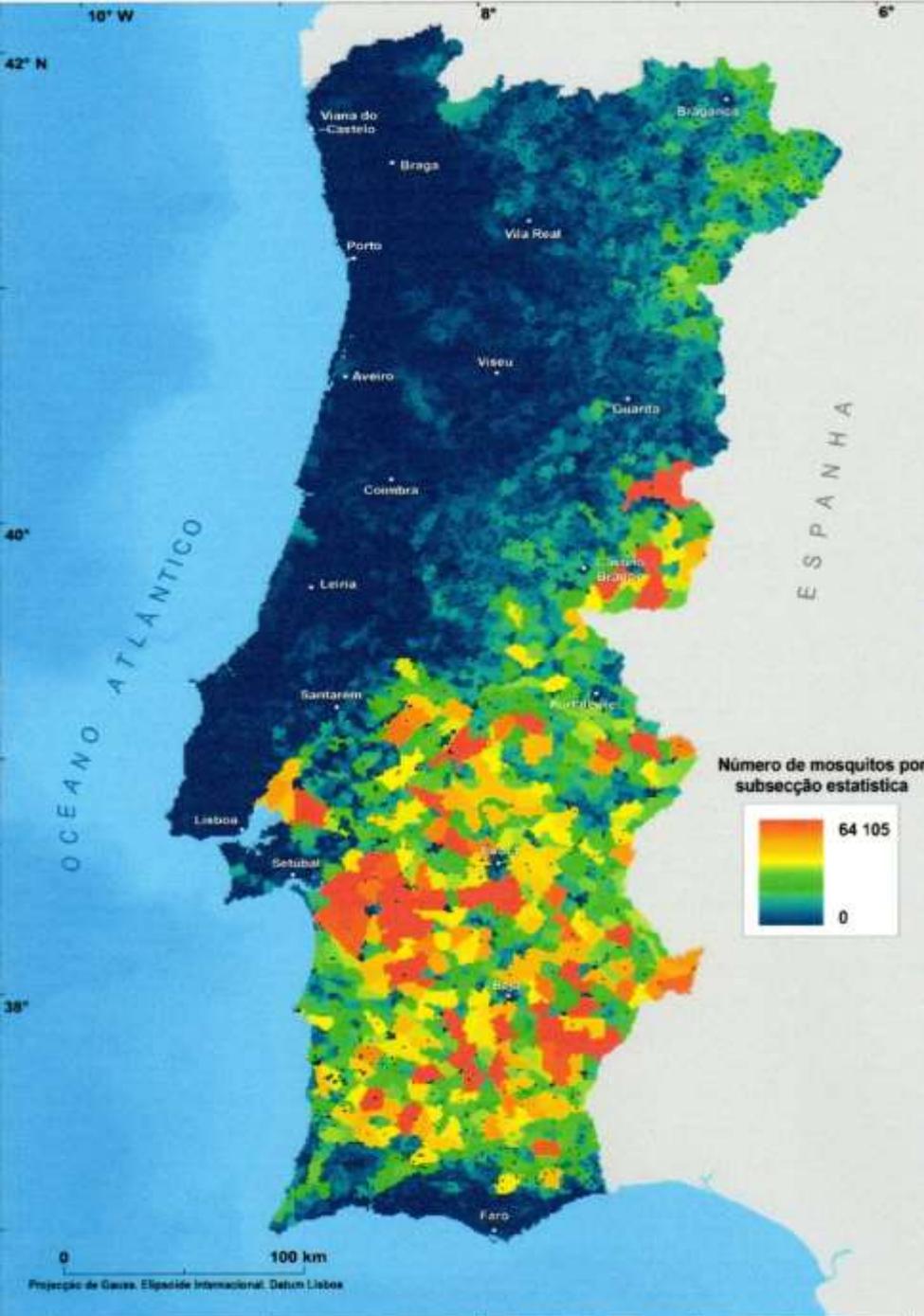


Figura 3.11 – Pessoas infectadas com malária (casos importados) em Portugal Continental, por concelho, no ano 2000 (fonte dos dados: DGS, 2001)



A.1 – Modelo preditivo de abundância de *Anopheles atroparvus* para Portugal Continental





A.3 – Número de *Anopheles atroparvus*, por subsecção estatística, em Portugal Continental

A.5 – Número de mosquitos por pessoa, por subsecção estatística, em Portugal Continental



# Conclusão ... “ ... não é completamente impossível que ... ”

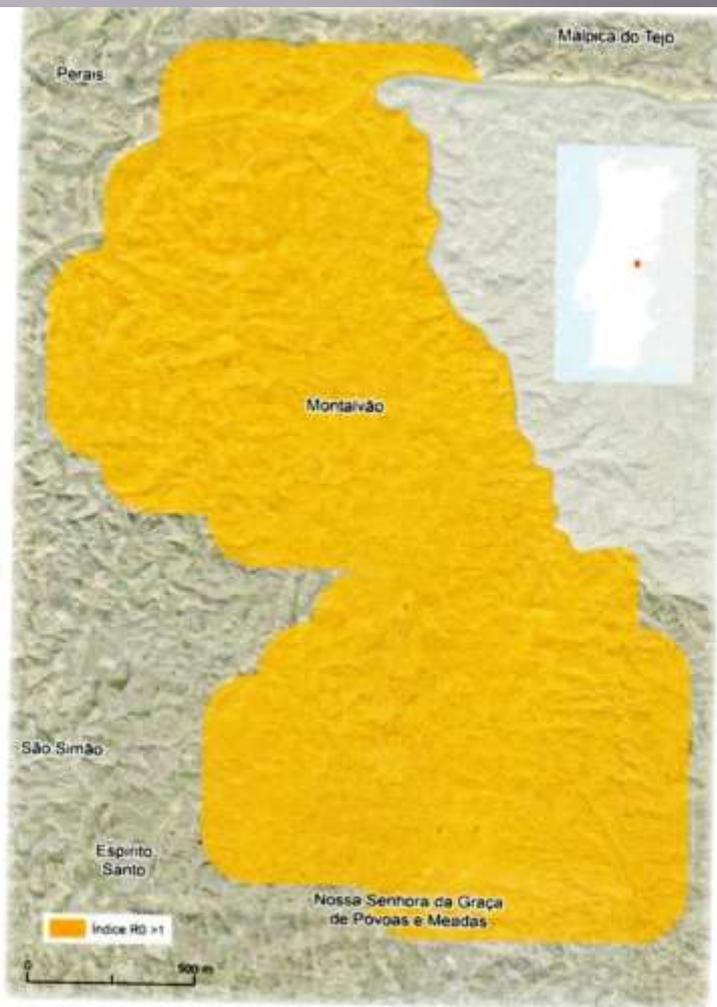


Figura 3.14 – Área de risco máximo de transmissão de malária (freguesia de Montalvão, concelho de Nisa)

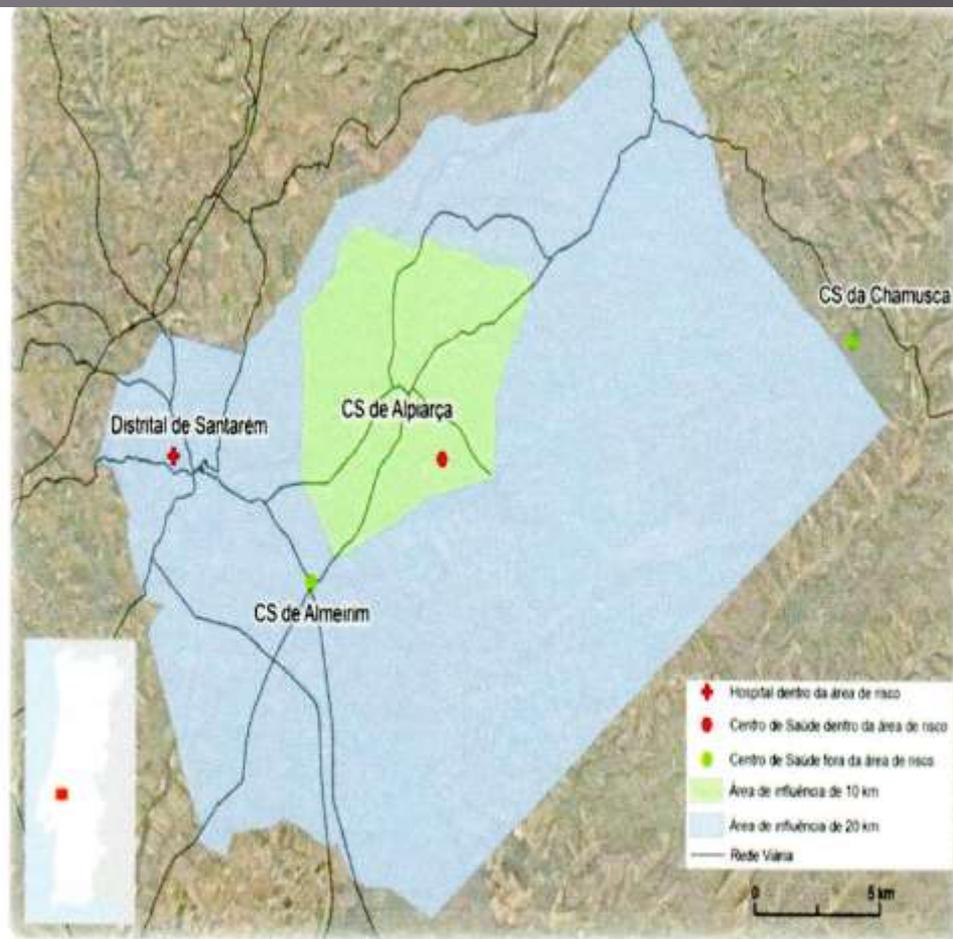
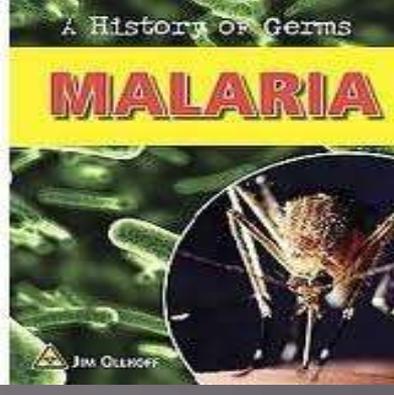
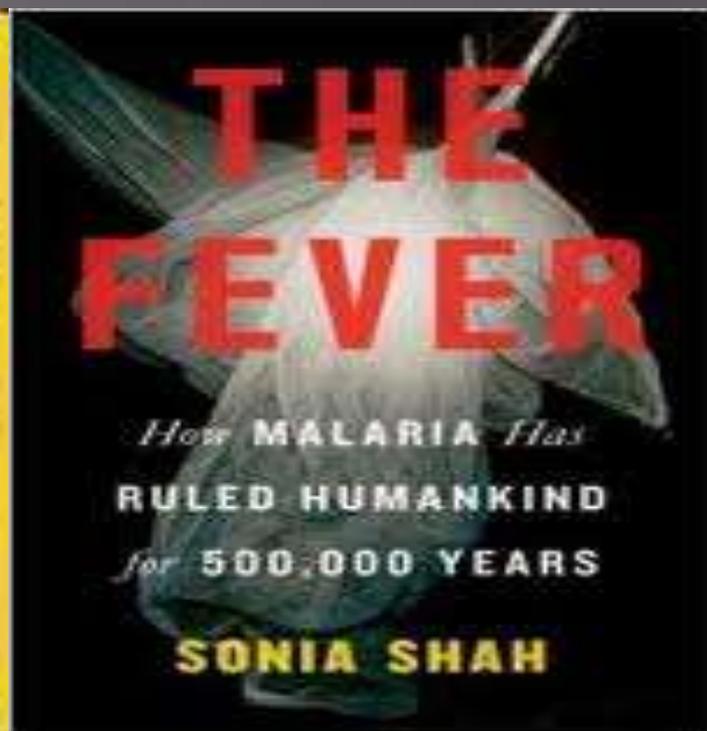
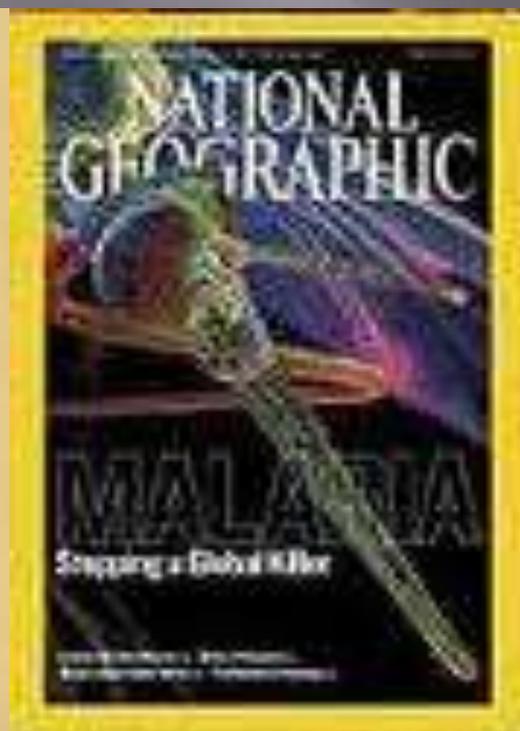


Figura 4.2 – Exemplo de identificação de hospitais e centros de saúde nas áreas de maior risco de transmissão de malária em Portugal Continental

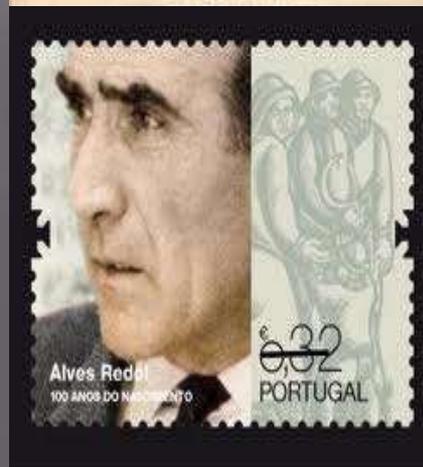
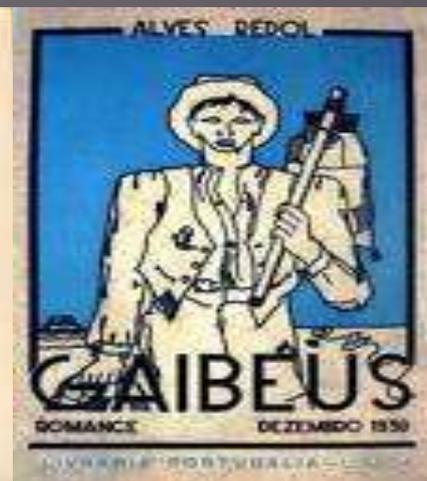
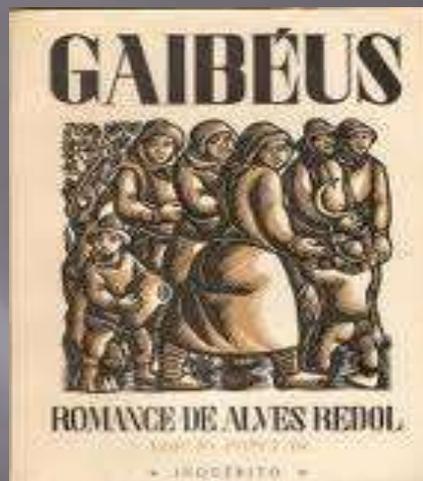


## CONTROVÉRSIAS E CURIOSIDADES DA HISTÓRIA: O CONTRIBUTO PORTUGUÊS





- *“ ... cada homem na eira não passa de um volante, uma correia ou um braço da ciranda. Quando o apito soar, o volante achará os raios, a correia e o braço da ciranda adormecerão. Os homens irão ajudar à carga e pensar na vida. Nos corpos de alguns correrá o frio das sezões; e os cérebros, libertos da vertigem comunicativa das máquinas, encontrarão pensamentos. Mas os pensamentos não sabem ainda acalentar fadigas. Nas poisadas, a vida torna-se mais negra... ”(sic.)*







# Applications actuelles de la malariathérapie\*

G. LUPASCU<sup>1</sup>

*Après un bref rappel historique et l'évocation des résultats obtenus dans les centres d'impaludation thérapeutique, dont certains ont eu une importance décisive pour l'étude du paludisme, l'auteur expose la situation actuelle et les perspectives en matière de malariathérapie.*

## Procedure and clinical assessments of malariotherapy: recent experience in 20 HIV patients

CHEN Xiaoping 陈小平, XIAO Binqun 肖斌权, XU Huifang 徐慧芳, SHI Wenjun 施文钧, GAO Kai 高凯 and RAO Jili 饶纪礼

**Keywords:** malariotherapy · human immunodeficiency virus · acquired immunodeficiency syndrome · side effect

**Objective** To demonstrate the side effects of malariotherapy and to explore safe procedures in conduct of malariotherapy for human immunodeficiency virus (HIV) infected patients.

**Methods** Twenty HIV/ acquired immunodeficiency syndrome (AIDS) patients were selected for the study of malariotherapy and were intravenously infected with Plasmodia vivax to induce therapeutic malaria. Malaria was terminated with chloroquine after 10 - 20 malarial febrile episodes. Clinical assessments were made before (baseline), during (malarial phase) and after (post) termination of malaria. The density of Plasmodia in peripheral blood from the HIV/AIDS patients were compared to that from HIV-negative naturally infected malarial patients who donated the blood for the therapeutically induced malaria. CD<sub>4</sub> cell baseline levels were correlated to the severity of malarial symptoms and parasitemia.

**Results** There were no significant differences of Plasmodium density between the HIV/AIDS patients injected with P. vivax and the HIV-negative blood donors. However, it was found that the HIV-positive patients had milder malarial symptoms and parasitemia with progressively lower CD<sub>4</sub> cell baseline levels. All patients developed every day or every other day fever episodes with headache and shaking chill. These symptoms were well tolerated with the aid of anti-pyretic medications. Spleen and liver enlargement were seen in 15 of 20 and 4 of 20 patients respectively. Transitory liver effects with increase of serum glutamic-pyruvic transaminase were seen in 2 of 20 during malarial phase. Most patients experienced mild to medium anemia and 6 of 20 patients developed thrombocytopenia during malarial phase. All these side effects disappeared after termination of malaria or within one month thereafter. No complications occurred in these patients.

**Conclusions** Therapeutically induced acute vivax malaria was well tolerated in 20 HIV-positive subjects who represented a range of CD<sub>4</sub> cell levels from 1868/ $\mu$ l to 15/ $\mu$ l. Malariotherapy did not induce complications while increasing CD<sub>4</sub> cell levels in most treated HIV/AIDS patients (results published elsewhere).

Malariotherapy and cancer.

MedLine Citation: PMID: 7010092 Owner: NLM Status: MEDLINE

**Abstract/OtherAbstract:** Malariotherapy merits a clinical trial as an adjuvant to conventional cancer therapy. This particular modality of treatment is a most potent stimulus of macrophage activity. These scavenger cells are widely believed to be an essential arm in the host's immune defenses against malignant disease, both as regards the processing of antigens and as killers of tumor cells. Significant too, malariotherapy, as formerly used to effectively treat some 16,000 patients with parietic neurosyphilis in one institution alone, before the advent of the penicillin age, has proved to be a particularly safe modality of treatment.

Authors: L B Greentree



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**MMWR**

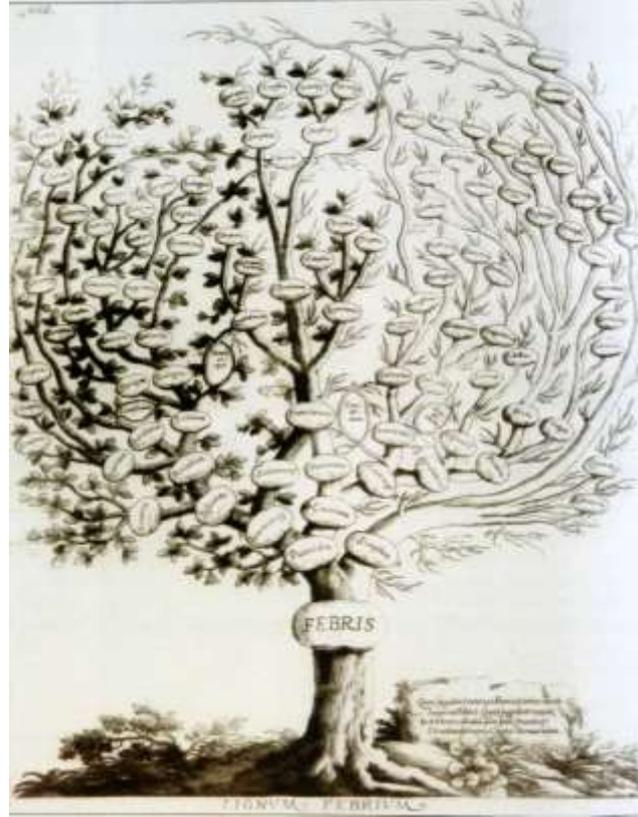
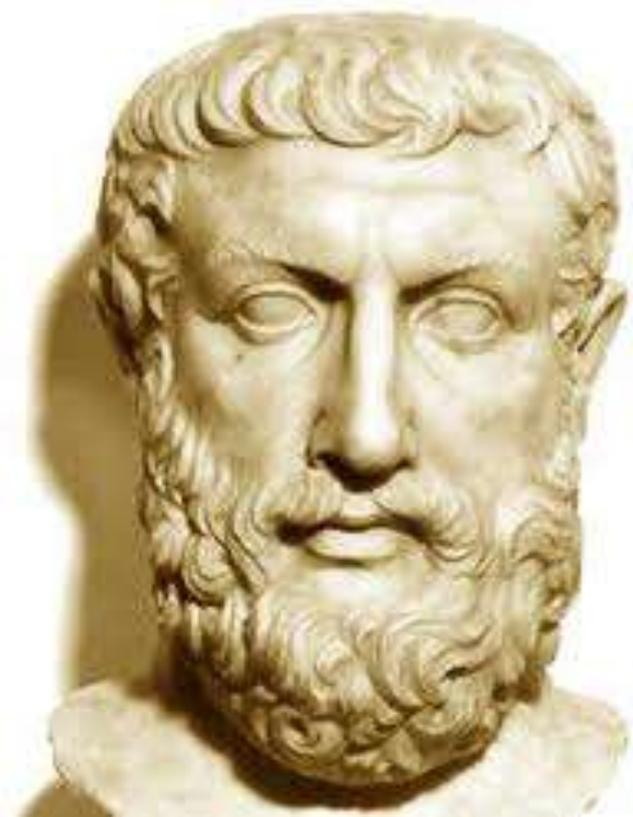
Weekly

October 04, 1991 / 40(39);665-666

## Update: Self-Induced Malaria Associated with Malariotherapy for Lyme Disease -- Texas

# A Febre como Arma Terapêutica I

- Parmenides (540-480 AC)
  - *"... give me the the power to induce fever, and I cure all the diseases..."*
- Thomas Sydenham (1624-1689)
  - *"... a Febre é um instrumento da Natureza para que o Homem possa combater os seus inimigos ..."*



# A Febre como Arma Terapêutica II

- Antoine Didier (1725)
  - Neoplasias em doentes c/ Sífilis tinham mais probabilidade de cura
  - Prostitutas tinham menor incidência de neoplasias
- W. Bucsh (1868)
  - Foi o 1º a utilizar a hipertermia induzida na terapêutica oncológica a partir de infecções provocadas p/ *S. pyogenes*
- Frederick Fehleisen (1882) e P. Bruns (1887)
  - Induziram remissão em vários doentes c/ neoplasias provocando iatrogenicamente uma erisipela p/ *S. pyogenes*
- E. Lambote (1896)
  - Constatou que os doentes que tinham tido erisipela mais raramente desenvolviam neoplasias
- William Cooley (1862-1936) (NY Hospital e Memorial Sloan-Kettering Cancer Center – USA)
  - Foi o 1º a estudar este fenómeno de forma sistemática através da utilização de uma Toxina (*Streptococos* e *Serratia*)
- P. Engel (1934-1935) e F. Sinek (1936)
  - Constataram que doentes que nunca tinham tido acessos febris tinha maior probabilidade de virem futuramente a desenvolver neoplasias
- R. Virchow (1863)
  - Estabeleceu a relação entre a infecção p/ *S. Haematobium* e o carcinoma da Bexiga
- M. Askanazy (1911)
  - Estabeleceu a relação entre a infecção p/ *O. Felineus* e o Carcinoma do Fígado

## Fever, Cancer Incidence and Spontaneous Remissions

Ralf Kleef<sup>a</sup> Wayne B. Jonas<sup>b</sup> Wolfgang Knogler<sup>c</sup> Werner Stenzinger<sup>d</sup>

<sup>a</sup>Office of Complementary and Alternative Medicine, NIH, <sup>b</sup>Samueli Institute for Information Biology and Uniformed Services University of the Health Sciences, Bethesda, Md, USA, <sup>c</sup>Menox Ambulatorien, Vienna, Austria, <sup>d</sup>Odenwaldklinik, Bad König, Germany

**Table 1.** References reporting infection and/or fever in association with spontaneous remission of neoplastic diseases

Tumor types	References
Bone tumors	112, 144-148
Burkitt's lymphoma	151
Brain tumors	122, 149-150
Colorectal cancer	123-124
Gastric cancer	152-153
Gynecological	154-155
Head and neck cancer	156-157
Hepatocellular cancer	158-161
Leukemia: AML, ALL, CML, CLL	115-117, 121, 125, 162-182
Lung cancer	119-120, 182-184
Lymphoma and non-Hodgkin lymphoma	151, 185-196
Melanoma	126, 197-202
Multiple myeloma	203
Prostate cancer	204
Renal cell cancer	205-207
Retinoblastoma	208-211
Sarcoma	95, 211-217

AML = Acute myeloic leukemia; ALL = acute lymphatic leukemia; CML = chronic myeloic leukemia; CLL = chronic lymphatic leukemia.

Uwe Hobohm

## Fever and cancer in perspective

Received: 22 March 2001 / Accepted: 5 July 2001 / Published online: 22 August 2001

Study	Treatment	Outcome	Remark	Ref.	Year
71 patients with inoperable metastatic neoplasms of different kinds, with different prior treatment by chemotherapy and radiation; 37 treated with CT vs. 34 with typhoid vaccine (control)	Coley's toxin	9 responses in CT, including 3 cures (no sign of residual tumor); 1 slight response in control	Doses were increased in both groups until 102°F was achieved; at least 10 injections were given; patients had different history of chemotherapy and radiation	[20]	1962
93 patients with inoperable metastatic neoplasms of different kinds, all treated by CT	Coley's toxin	18 responses including 1 cure	Doses were increased in both groups until 102°F was achieved; at least 10 injections were given; patients had different history of chemotherapy and radiation	[21]	1962

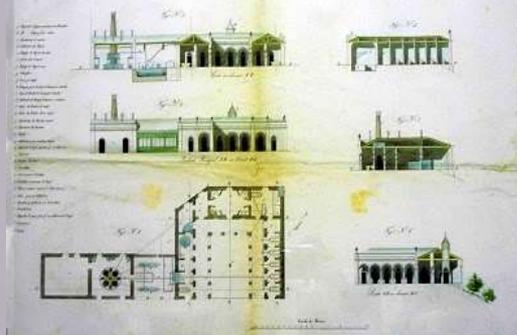


State of Nevada }  
Humboldt County }

On this the 13<sup>th</sup> day of June A.D. 1863  
Was Personally appeared before me E. A. Clark Judge  
of Probate in and for the aforesaid County and  
State Donald McKay one of the Witnesses of the  
Last Will and Testament of William Cooley  
deceased Who under oath says that the foregoing  
a Paper exhibited by Francis Matthews Son and  
of the deceased is the true and only last Will  
and Testament of the said William Cooley and  
Donald McKay  
Saw me to and subscribed before me  
this 13<sup>th</sup> day of June A.D. 1863  
E. A. Clark  
Judge of Probate



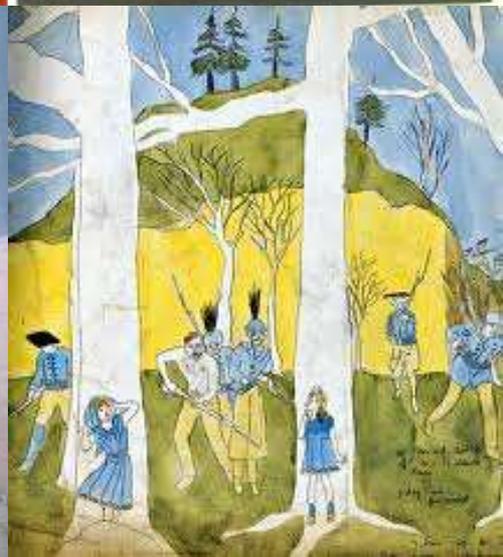
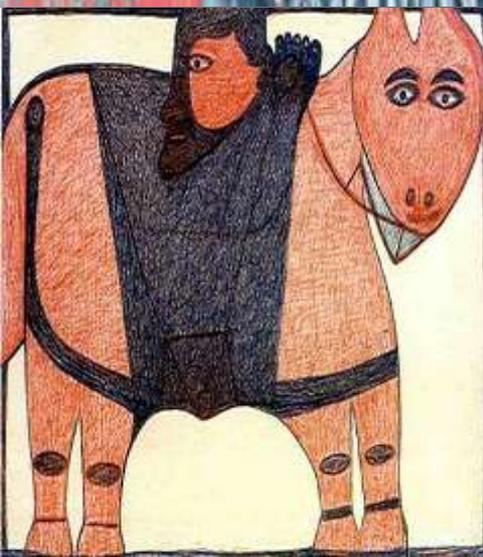
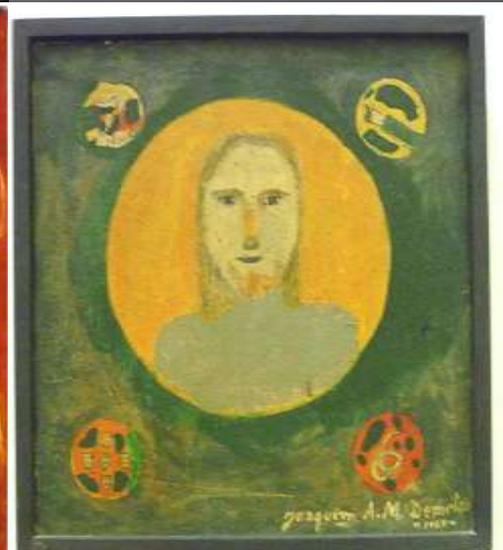




GATAFUNHOS  
E  
OUTRAS ARTES

Projeto de Arte em  
Apoio à  
Saúde Mental

HOSPITAL TIQUEL SOBRADA,  
HOSPITAL Nº 204 - 15 Março 2005  
Rua Santa Fátima, 161  
HORAIS 10H às 15H



## 4.2. Malarioterapia

Operação que visa tratar e curar enfermidades nervosas por introdução do *plasmodium vivax*, do hematozoário de Laveran no organismo humano. Desde 1917 que a malarioterapia foi aplicada em paráliticos gerais e associada ao tratamento da sífilis precoce. Foi o psiquiatra austríaco Julius Warner-Jauregg (1857-1949) que descobriu a importância terapêutica da inoculação de malária no tratamento da paralisia demencial, o que lhe valeu o Prémio Nobel da Medicina em 1927.

Em Águas de Moura realizaram-se trabalhos de malária experimental em articulação com o Hospital Miguel Bombarda. Estas experiências decorreram a partir de 1937, destinados ao tratamento de casos de sífilis cerebral. A documentação existente refere-se ao período 1937-1942.

Em 1953 há ainda referência ao prosseguimento de malarização com espécimes do Instituto.

1. A malarioterapia na cura da parálisia geral sífilítica / J. Menezes de Almeida. Porto: Lopes da Silva - Editora, 1950.

Adicionar à lista

2. Luta anti-malária e malarioterapia: relatório duma viagem de estudo: 1932 / José Chaves Ferreira. Francisco Cambournac. [S.l. : s.n.], 1935.

Adicionar à lista

3. Nota de um clínico geral sobre a malarioterapia / por J. Gouveia Monteiro. [S.l. : s.n.], 1954.

Adicionar à lista

4. XXV Concurso Hípico Internacional Oficial. [S.l. : s.n.], 1938.

Adicionar à lista

## Parasitological and Clinical Investigations on Infections with the VS Romanian Strain of *Plasmodium malariae* Transmitted by *Anopheles labranchiae atroparvus*\*

G. LUPASCU,<sup>1</sup> P. CONSTANTINESCU,<sup>2</sup> E. NEGULICI,<sup>3</sup> P. G. SHUTE<sup>4</sup>  
& M. E. MARYON<sup>5</sup>

*In order to contribute to knowledge of the sporogonic cycle of malaria parasites, 3 batches of Anopheles labranchiae atroparvus were infected with the VS Romanian strain of Plasmodium malariae by feeding them on 2 patients undergoing malariotherapy in Romania. Of the 310 infected mosquitos 59 were dissected for ookinetes, oocysts and sporozoites, and the rest were used for further clinical trials in England. Mature ookinetes were observed 18 hours after infection, and sporozoites were first observed in the salivary glands 19 days after infection.*

*One of the main characteristics of the VS strain of P. malariae observed in Romania was the relatively large number of gametocytes and subsequent heavy infections it could produce. However, gametocytes were found only in the heart blood of 1 of the 4 subjects infected in England (on autopsy after accidental death); post mortem examination of the liver of the same subject failed to show exo-erythrocytic schizonts. None of the other 3 subjects showed gametocytes or infected mosquitos. On the basis of clinical symptoms, the incubation period of P. malariae in these subjects was found to be 18-19 days.*

## FRANCISCO CAMBOURNAC



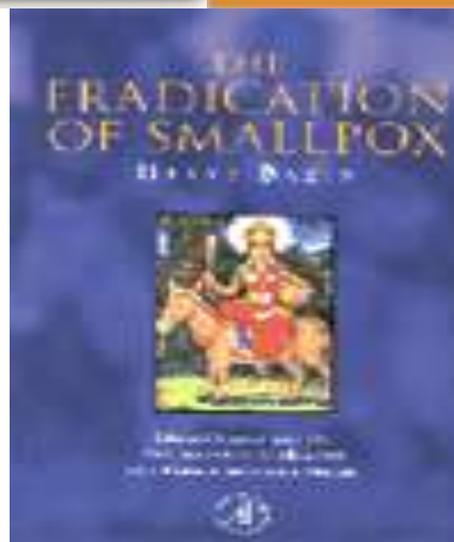
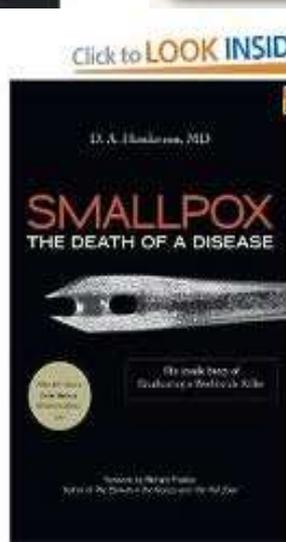
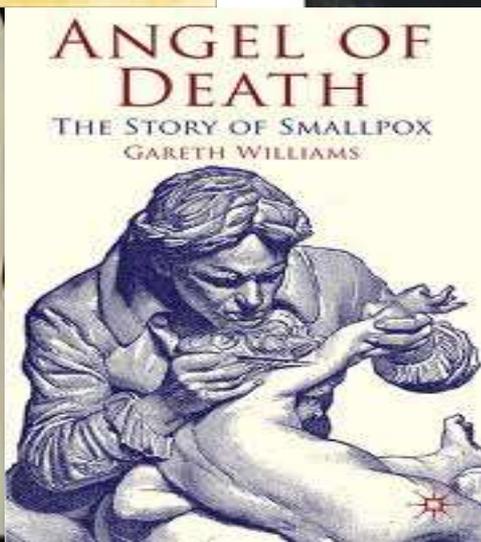
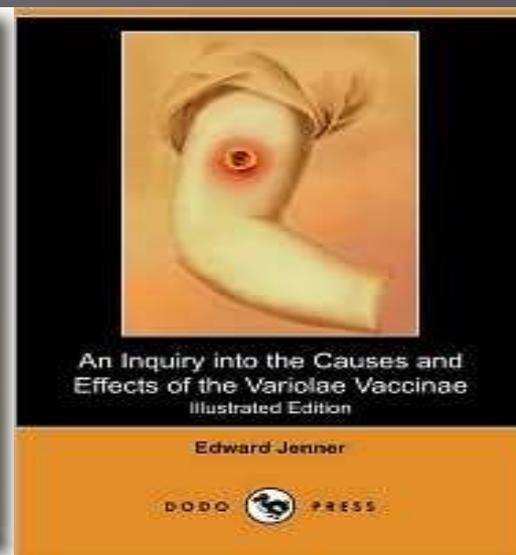
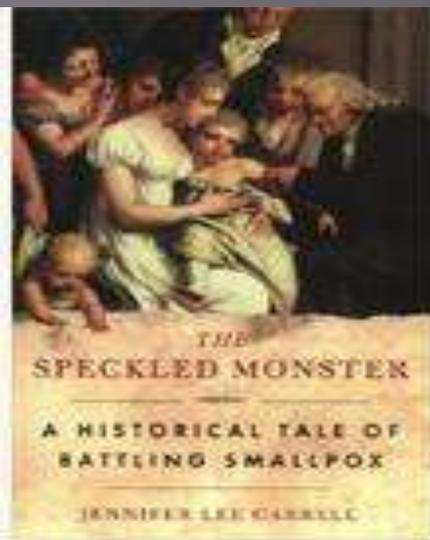
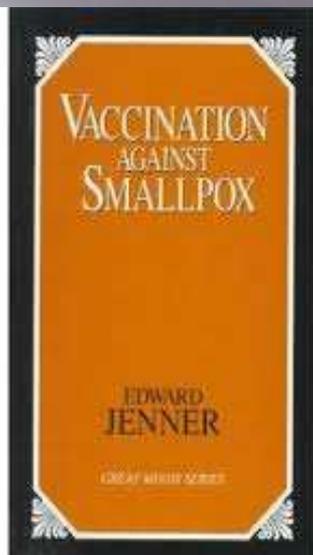
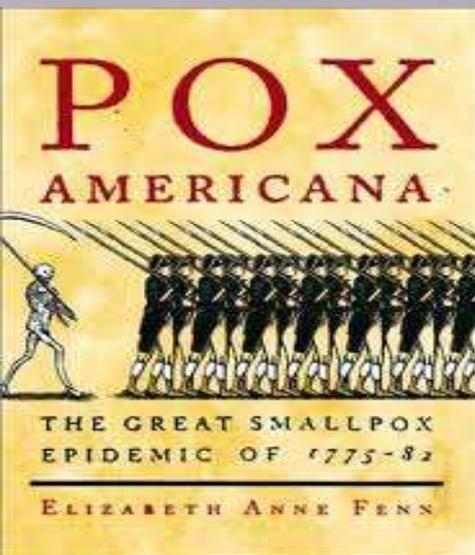
Médico epidemiologista e tropicalista, Francisco Jo Cambournac (1903-1994) destacou-se sobretudo Malariaologia, área em que deu um grande contrib portuguesa. Além de membro-fundador e director Malariaologia de Águas de Moura (1939-1954), Francisco ainda director da Organização Mundial da Saúde para durante dez anos (OMS África), tendo recebido o Prém (1978).

Licenciado em Medicina pela Faculdade de Medicina da Lisboa, Francisco Cambournac fez a sua formação hosj sua carreira clínica no Hospital de Santa Marta, er posteriormente concluído o curso da antiga Escola de Me

Decidido a especializar-se em doenças tropicais, em particular na área de Malariolc curso de Malariaologia da Faculdade de Medicina de Paris, fez estágios prático Jugoslávia, nomeadamente na Escola Superior de Malariaologia e na Estação Exp luta anti-malária de Roma, e no Instituto de Higiene de Skoplje. Frequentou a Higiene e Medicina Tropical no Instituto de Medicina Tropical de Hamburgo e rea Instituto Pasteur de Paris, no Instituto Colonial de Amesterdão, na Escola de Hi Tropical de Londres, no Ross Institute e na Secção de Malarioterapia do Horton M Epsom.



# Variola: A 1ª Doença Erradicada pela Vacinação (História de um Contributo Esquecido)



# La croisade antivariolique

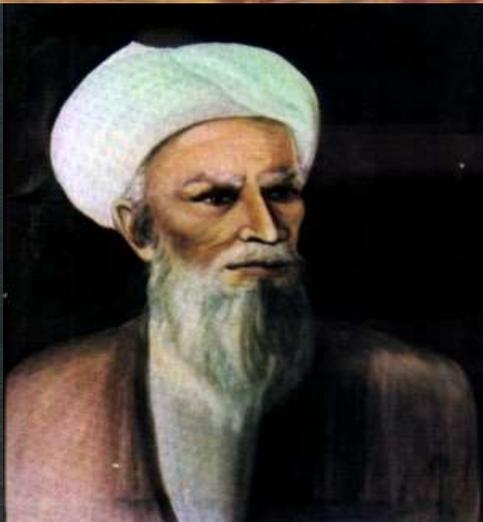
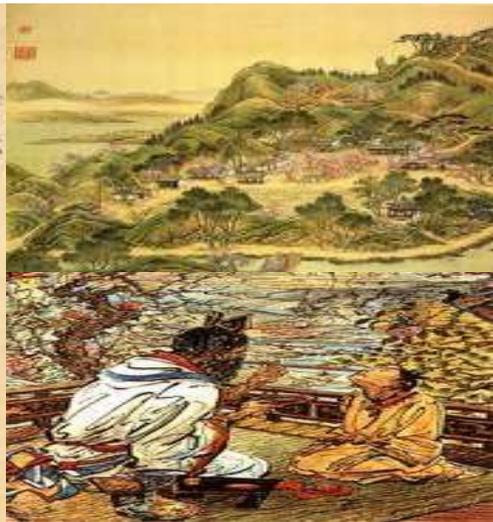
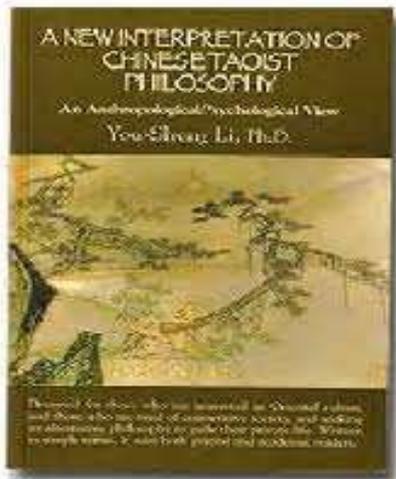
par Pierre Darmon

## Il a fallu un siècle pour vaincre les craintes des Français devant une vaccination antivariolique qui n'était d'ailleurs pas sans risques.

Table 6.1. Important events in the history of smallpox control, from ancient times to 1900

Variolation	Vaccination	Isolation and quarantine
<b>10th century</b> Variolation first reported in China, by insufflation, as a secret rite. Probably also practised in India at this time, by cutaneous inoculation.		Hospitals for smallpox established in Japan (Ashino, 982).
<b>13th century</b> Variolation by cutaneous route introduced into Egypt by Mamelukes.		
<b>17th century</b> Variolation more widely used in China. Kang Hsi (1661-1722) variolated his soldiers and his children.		Quarantine introduced to control entry of smallpox into North American ports (Boston, New York, Philadelphia: 1650s). Mandatory isolation of smallpox cases at home (Virginia: 1667).
<b>18th century</b> Papers on variolation published by Royal Society of London (Chinese method, 1700; Turkish method: Timoni, 1714; Fyhririni, 1716). Cotton Mather told of variolation by his African slaves (Boston, 1706). Variolation by cutaneous route carried out in Great Britain (Sloane, 1721), Bohemia (Reiman, 1721) and Boston, USA (Boylston, 1721). Variolation popularized in England by the Suttons (1726). Dimsdale variolates Catherine the Great and variolation accepted in Russia (1748). Louis XV dies of smallpox and variolation accepted in France (1774). Washington orders variolation of the Continental army (1777).	Publication of Jenner's inquiry (1798).	London Small-Pox and Inoculation Hospital established (1746). Eradication of smallpox by systematic variolation of population and isolation of cases suggested by Haygarth (1793) and Carl (1799).
<b>19th century</b> Variolation banned in Russia (1805), Prussia (1835), Great Britain (1840) and British India (1870), but still widely practised in Afghanistan, China and many parts of Africa.	<i>Inquiry</i> translated into several European languages (1800-1802). Vaccination adopted in most European countries and in the USA (1800-1803). Vaccine sent successfully to Bombay (de Carro, 1802) and to South and Central America, the Philippines and Macao (Balnis-Salvany Expedition, 1803-1806). Primary vaccination of infants made compulsory in Bavaria (1807), Denmark (1810), Norway (1811), Bohemia and Prussia (1812), Sweden (1816), Hanover (1821) and Great Britain (1853). Revaccination introduced into Württemberg (1829). Vaccination compulsory in Prussian army (1833). Vaccine produced in calves (Italy, 1805, 1810). Vaccine passaged in calves for production (Negri, 1840). Production in calves adopted in France (1864), Belgium (1865), Great Britain (1881) and Germany (1884). Use of glycerol as diluent introduced in Italy (Negri, 1840s). Glycerinated vaccine popularized by Coppenhan (1892). Jenner's arm-to-arm vaccination banned in Great Britain (1898).	Control of smallpox by isolation of cases and quarantine of contacts ("Leicester method", 1870); reinforced by vaccination of contacts (Billard, 1914).

# الحياة والصحة الأبدية: الطب الصيني



# The Montague Millennium

1000 years of worldwide family history



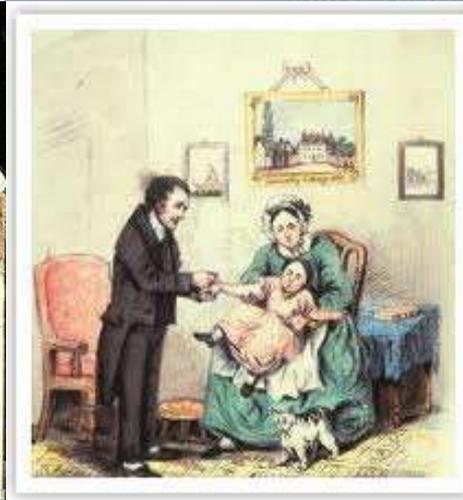
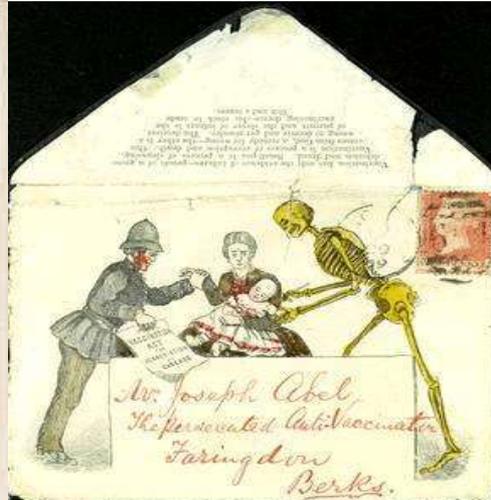
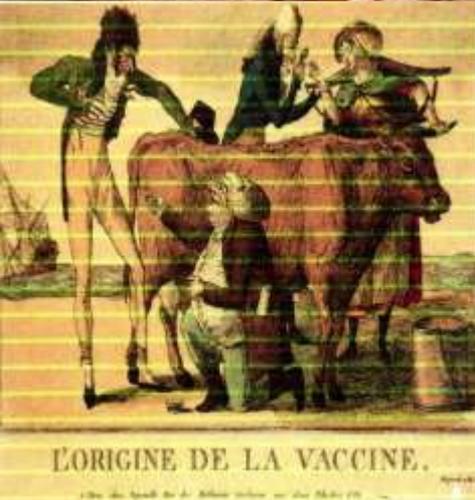
A smallpox vaccine commemoration stamp (Turkey: 1967)

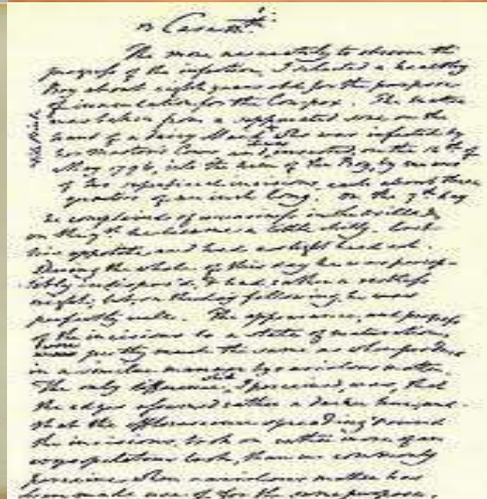
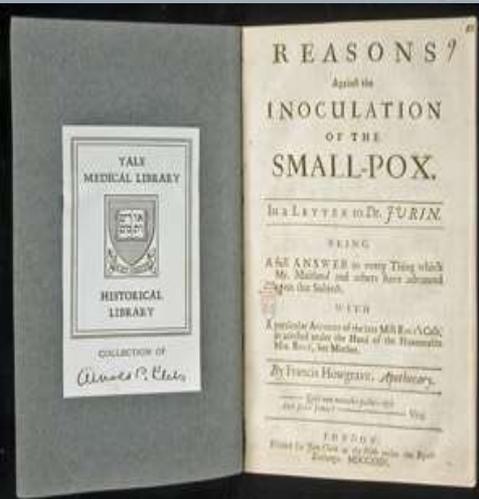
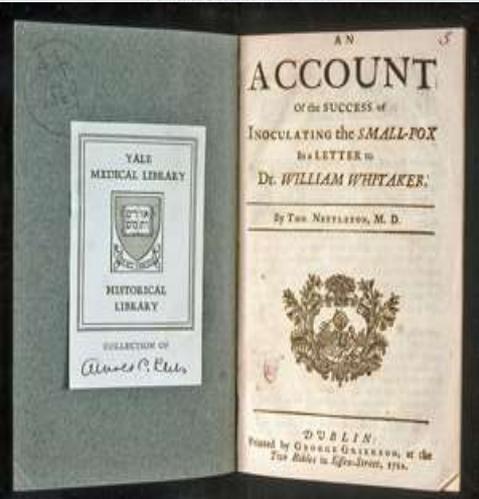
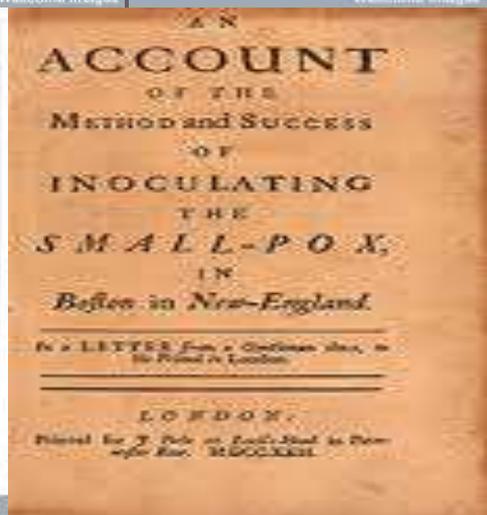
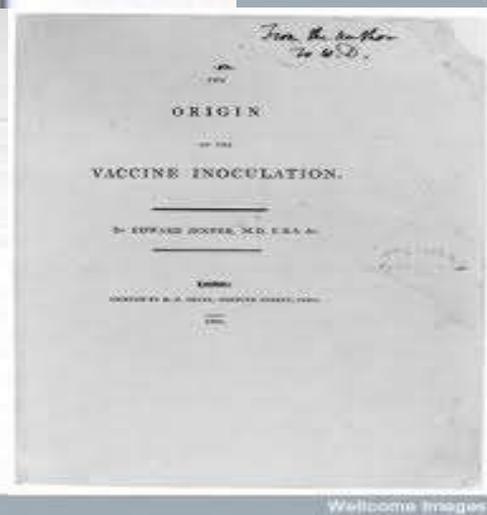
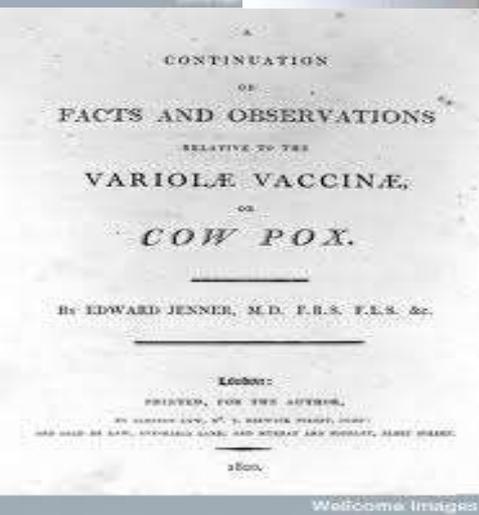
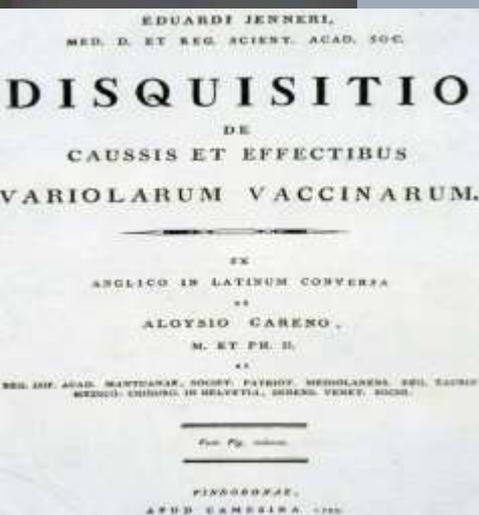
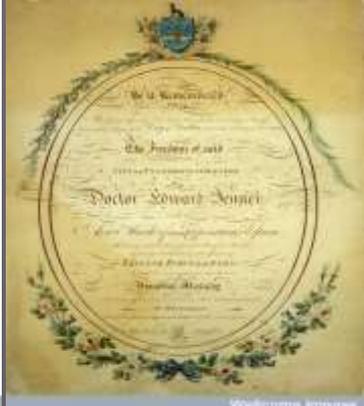
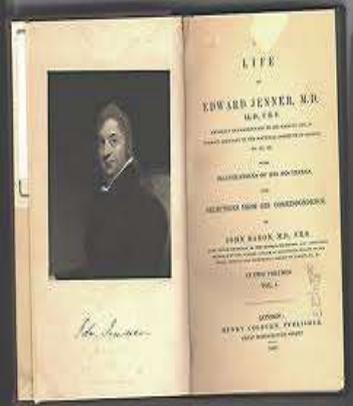






Welcome-Images

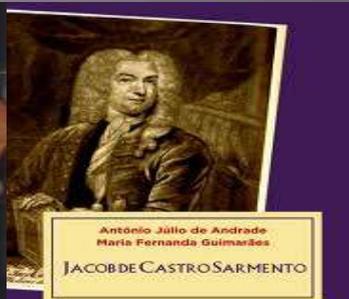
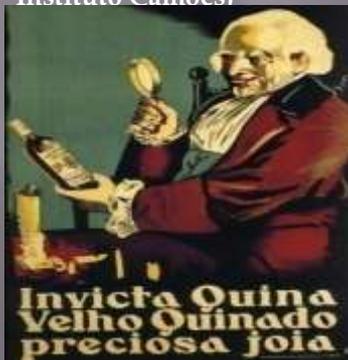


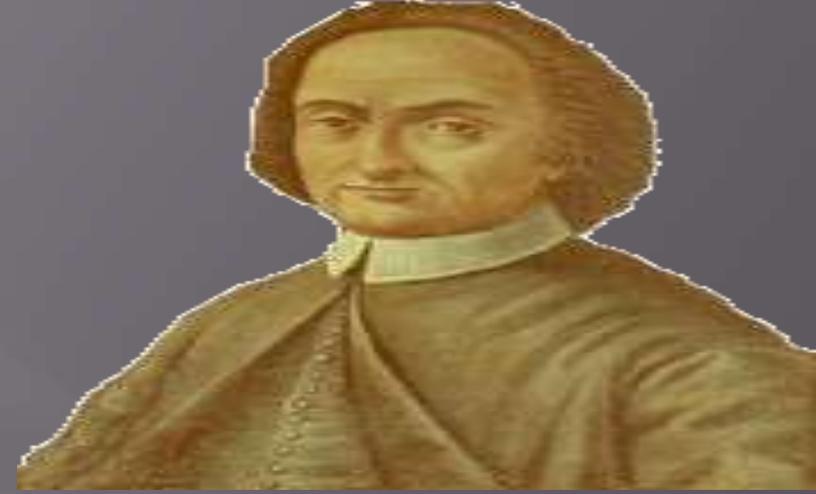




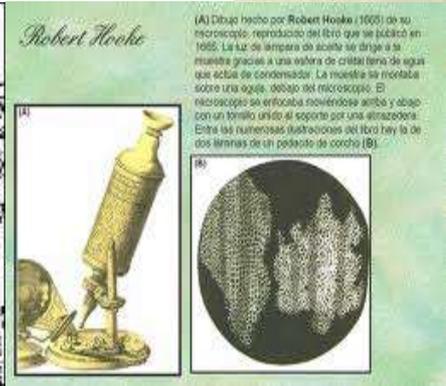
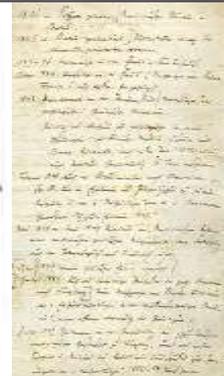
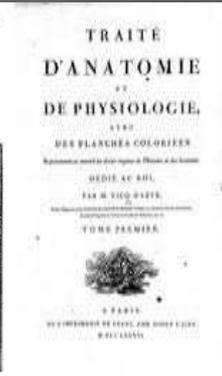
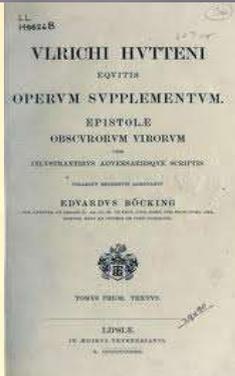
# A Água de Inglaterra

- “Água de Inglaterra é um dos exemplos mais marcantes dos 'remédios de segredo' muito em voga durante o século XVIII. Pelo nome de Água de Inglaterra eram conhecidos vários preparados farmacêuticos, produzidos por diferentes fabricantes desde finais do Séc. XVII a inícios do Séc. XIX e que apresentam em comum, além do nome, o facto de serem vinhos de quina. Eram utilizados para o tratamento do paludismo, que era então uma das doenças mais importantes, atingindo uma grande número de indivíduos e existindo endemicamente em várias regiões de Portugal. A importância medicinal da Água de Inglaterra reside principalmente no seu efectivo valor terapêutico, pelo facto de a quinina ser o seu princípio activo mais importante, constituindo o mais antigo quimioterápico ainda em uso. Era um medicamento popular, amplamente conhecido e divulgado, sendo frequentemente consumido por automedicação. Numa primeira fase era importada de Inglaterra, de onde Fernando Mendes (?-1724), o seu introdutor em Portugal, a enviava. Seguiu-se-lhe Castro Sarmento que montou uma verdadeira rede de distribuição da Água de Inglaterra em Portugal. A pouco e pouco foram surgindo produtores locais que foram aumentando a produção de forma a satisfazer a procura. Com o isolamento da quinina por Pelletier (1788-1842) e Caventou (1795-1877) em 1820 é a sua substituição pelo sulfato de quinina, perde a importância que teve no século XVIII.” (in Instituto Camões)



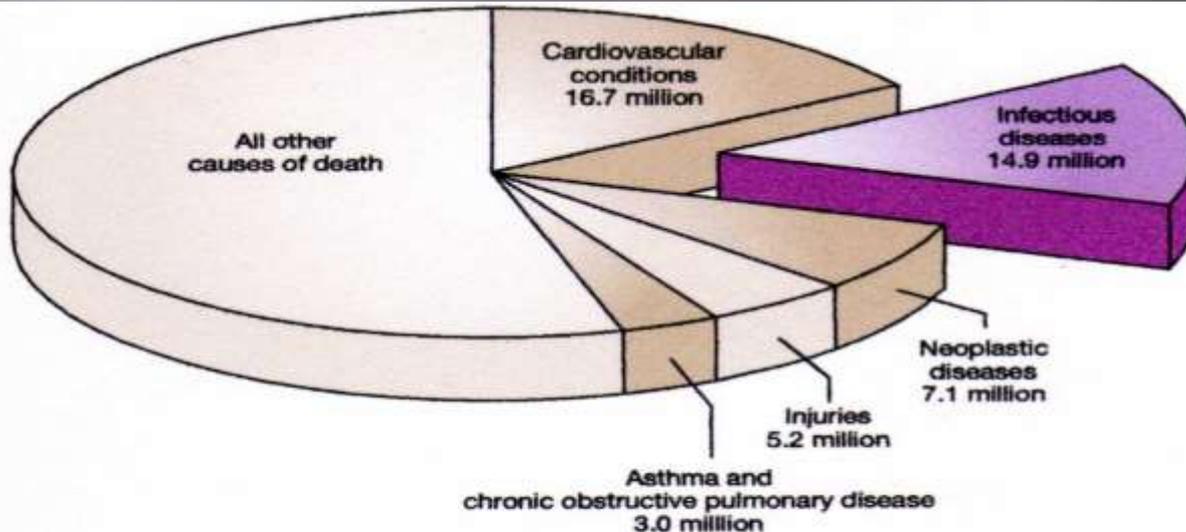
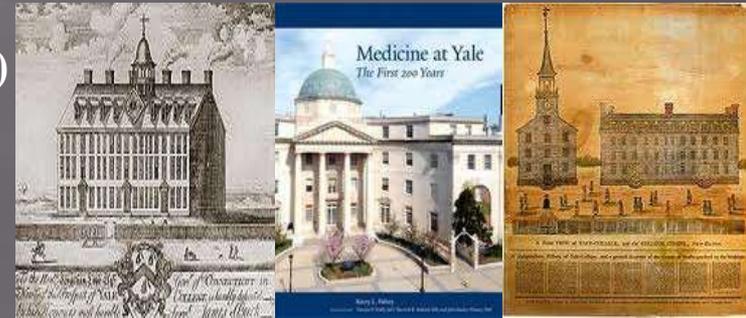


# (ALGUMAS) CONCLUSÕES



# Futurologia Falaciosa

- "... the war against infectious diseases is over, because we have won..." (sic.) (US Surgeon General, 1967)
- "... no new diseases are going to be discovered ..." (sic.) L. Thomas Dean of Yale Medical School (1976)
- WHO (2008)
  - D. Infecciosas 15.000.000 mortes / ano
    - 2ª causa de Mortalidade Mundial
      - 1ª Causa nos Países em Desenvolvimento
  - Novos Agentes Patogénicos para a Espécie Humanas descritos nos últimos 40 anos: 50



Infectious diseases	Annual deaths (million)
Respiratory infections	3.96
HIV/AIDS	2.77
Diarrhoeal diseases	1.80
Tuberculosis	1.56
Vaccine-preventable childhood diseases	1.12
Malaria	1.27
STDs (other than HIV)	0.18
Meningitis	0.17
Hepatitis B and C	0.16
Tropical parasitic diseases	0.13
Dengue	0.02
Other infectious diseases	1.76

**Figure 2** Leading causes of death worldwide. About 15 million (>25%) of 57 million annual deaths worldwide are the direct result of infectious disease. Figures published by the World Health Organization (see <http://www.who.int/whr/en> and ref. 7).



“...nowadays we see maladies unknown to our forefathers springing up around us”.

Ullrich von Hutten, 1519<sup>36</sup>



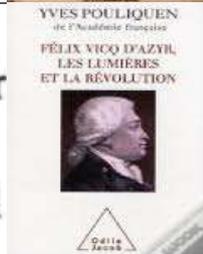
“After our fathers and grandfathers succumbed... dogs and vultures devoured the bodies. So it was that we became orphans, oh, my sons! So we became when we were young. All of us were thus. We were born to die!”

Cakchiquel Mayan on the huey zahuatl epidemic of 1520–21<sup>44</sup>



“If there is, within medicine, any subject worthy of... investigation... it is, without doubt, epidemic pestilential diseases, obscure and hidden in their causes, rapid in their progression, frightening in their symptoms, and deadly in their consequences.”

Félix Vicq-d’Azyr, 1776<sup>51</sup>



“France and its capital have been visited by a fearful pestilence... It comes without any known cause; it disappears without any revealed reason. The bodies of its victims are in vain examined; death is interrogated: death betrays nothing.”

Louis-François Benoiston de Châteauneuf, 1834<sup>69</sup>



“Not a single year passes without [which]... we can tell the world: here is a new disease!”

Rudolf Virchow, 1867<sup>1</sup>



*“The future of microbes and mankind will probably unfold as episodes of a suspense thriller that could be entitled Our wits versus their genes”.*

J. Lederberg, *Science* 2000<sup>33</sup>



Joshua Lederberg (1925 - 2008)

## Charles Nicolle, Prémio Nóbel da Medicina, 1933

- “... irão haver sempre novas doenças. É um fatalidade. Uma outra fatalidade, é o facto de nós não sermos imediatamente capazes de determinar a sua etiologia. Devemo-nos pois resignar com a ignorância perante os primeiros casos. Eles serão, pelo desconhecimento, encarados como uma doença já conhecida ...” sic.



## Portugueses desenvolvem vacina contra a malária

Fundação Melinda & Bill Gates financia pela primeira vez projetos lusos, que são pioneiros a nível mundial.

# INSTITUTO GULBENKIAN de CIÊNCIA



Comissário de Honra  
**Armindo R. Filipe - Director do CEVDI**

Organização  
**Câmara Municipal de Palmela**  
**Departamento de Cultura e Desporto**  
**Divisão de Património Cultural - Museu Municipal**

Colaboração  
**Câmara Municipal do Montijo**

Programação e Textos da Exposição  
**Ireneu Cruz**  
**Isabel Lucas**  
**Maria Teresa Rosendo**

Inventariação e Seleção de espólio  
**Isabel Lucas e Maria Teresa Rosendo**



A realização de um Simposium de homenagem aos Prof. Doutores Armindo Filipe e David Morais sobre a problemática actual e futura das Doenças Infecciosas, não faria sentido sem em simultâneo, chamar a devida atenção da sua importância histórica enquanto ex-Instituto de Malariologia e, numa derradeira tentativa, tentar contribuir para salvar o seu valiosíssimo património museológico.

A erradicação da malária no território nacional, foi uma epopeia de enorme envergadura, que encerra seguramente motivos importantes de reflexão para os responsáveis pela nossa política de saúde, dado que, não só se podem aplicar a outras (novas e velhas) doenças transmissíveis, mas também porque, segundo a previsão de alguns credenciados cientistas, esta doença poderá voltar a afectar de novo os países da orla mediterrânica.

Já é tempo de interiorizarmos definitivamente a ideia que o Fim das Doenças Infecciosas é uma mera quimera sem qualquer fundamento científico credível, e que as Doenças Emergentes, Reemergentes e de Importação constituem e constituirão cada vez mais uma ameaça quotidiana à saúde das populações de todo o nosso planeta.

Só compreendendo as lições da nossa história colectiva e respeitando os seus verdadeiros protagonistas poderemos ambicionar estar à altura destes enormes desafios.

O objectivo principal deste Simposium, bem como da realização da Exposição e da edição do seu Catálogo é pois servir esta nobilíssima causa.

O Responsável da Comissão Organizadora do Simposium  
**José Poças**



## Simposium Doenças Infecciosas – Perspectivas no Virar do Milénio Cine-Teatro S. João - Palmela, 28 de Novembro a 2 de Dezembro de 2001

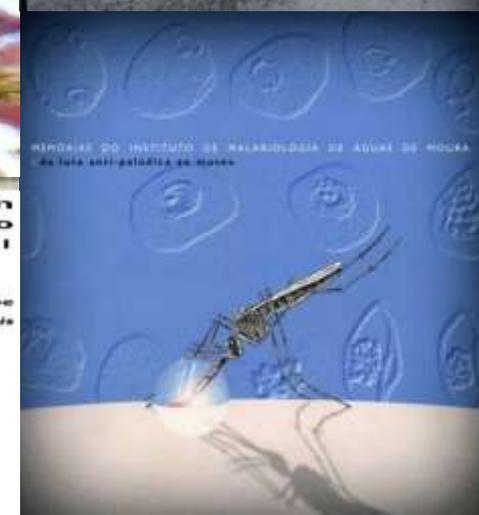
Presidentes: **Prof. Doutor Armindo Filipe**  
 Presidente Honorário: **Prof. Doutor David Morais**

### Organização

Hospital de S. Bernardo – Valência de Doenças Infecciosas  
 Instituto Nacional de Saúde Dr. Ricardo Jorge – Centro de Estudos de Vectores e Doenças Infecciosas

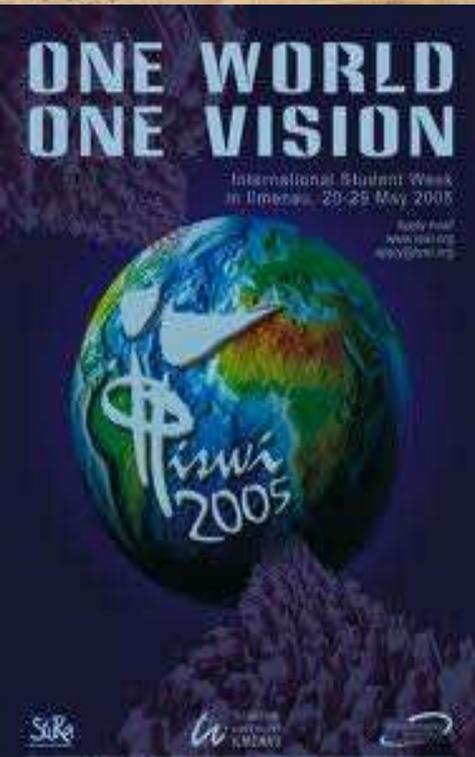
### Comissão Organizadora

José Poças (Responsável), Ireneu Cruz, Isabel Lucas, Maria Teresa Rosendo, José Barata,  
 Eduardo Carqueijeiro, António Xavier, Fátima Bacellar, Joana Sá, Maria João Alves, Nathalie Graupner,  
 Paula Proença, Sofia Nuncio





THE LANCET Infectious Diseases



## Towards a conceptual framework to support one-health research for policy on emerging zoonoses

Richard Coker, Jonathan Rushton, Sandra Mounier-Jack, Eshan Karimuribo, Pascoal Lutumba, Domsinic Kombarage, Dirk U Pfeiffer, Katharina Stark, Mark Rweyemamu

In the past two decades there has been a growing realisation that the livestock sector was in a process of change, resulting from an expansion of intensive animal production systems and trade to meet a globalised world's increasing demand for livestock products. One unintended consequence has been the emergence and spread of transboundary animal diseases and, more specifically, the resurgence and emergence of zoonotic diseases. Concurrent with changes in the livestock sector, contact with wildlife has increased. This development has increased the risk of transmission of infections from wildlife to human beings and livestock. Two overarching questions arise with respect to the real and perceived threat from emerging infectious diseases: why are these problems arising with increasing frequency, and how should we manage and control them? A clear conceptual research framework can provide a guide to ensure a research strategy that coherently links to the overarching goals of policy makers. We propose such a new framework in support of a research and policy-generation strategy to help to address the challenges posed by emerging zoonoses.

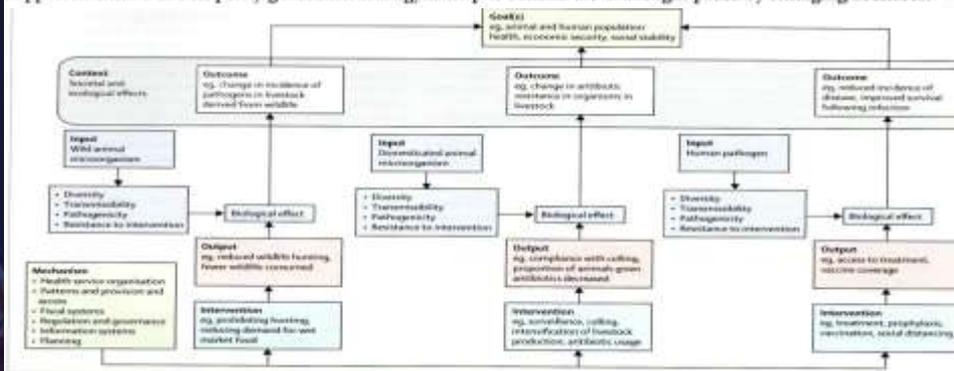


Figure: Schematic representation of a Framework for research to inform one health policy

STEPHEN MICHAEL APATOW

SM AEMDA COMMUNICATIONS: INTERNATIONAL

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