

Original Article

Surveillance of travel-associated diseases at two referral centres in Marseille, France: a 12-year survey

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Abstract

Background: With increasing international travel and historically high numbers of residents visiting friends and relatives overseas, travel-associated illnesses are frequent in Marseille, France. We report the changing epidemiology of travel-related illnesses over a 12-year period.

Methods: A single site GeoSentinel surveillance analysis was undertaken for 3460 ill returned travellers presenting to two public hospitals in Marseille, France from March 2003 to October 2015, with travel-related illnesses. Demographic characteristics, travel history, presenting symptoms and information on pre-travel consultations were collected.

Results: There was a predominance of travel to sub-Saharan Africa, in particular to Comoros archipelago. Tourism was the main reason for travel (1591/3460, 46%), followed by visiting friends or relatives (VFR) (895/3460, 26%), with a mean duration of 29 days; 35% (1212/3460) of travellers reported a pre-travel health consultation. The most common syndromic diagnoses were febrile systemic illness (1343, 39%), dermatologic (716, 21%), gastrointestinal (340, 10%) and respiratory/ear–nose–throat (331, ENT) (10%). Hospitalization rates were highest amongst travellers from sub-Saharan Africa (858/ 1632, 53%), and VFR (573/ 895, 64%, $P < 0.001$). Frequent diagnoses included malaria (797, 23%), dengue (96, 2.77%) and chikungunya (75, 2.17%), reflecting global trends. Comparison of two periods (2003–10 to 2011–15) demonstrated an increase in chikungunya and decrease in malaria and influenza-like illness. We report an increase in ill travellers from the Caribbean, Middle East and South-East Asia.

Conclusion: Surveillance of travellers provides relevant sentinel information on the changing epidemiology of infectious diseases across the globe, most notably for malaria, dengue and chikungunya. We demonstrate the use of travel surveillance in improving pre-travel consultation needs and to address autochthonous vector-borne viral risks.

Key words: Travel medicine, sentinel surveillance, malaria

Background

International travel is increasing, with more than 23 million visits overseas by French tourists in 2015, compared to over 84 million international tourists arriving in France over the same period.¹ Marseille, the main town in Mediterranean France, includes an urban and peri-urban population of over 1.5 million inhabitants. The economy of Marseille and its region is still partly linked to its commercial port and has always been one of

the main gateways into France.² Migrants, originating notably from North Africa, Senegal and Comoros, have settled in Marseille over several decades.^{2,3} Considering this context, travel-associated diseases are frequent in the local population.

We aim to report surveillance data to document the spectrum of infections in patients admitted for travel-related illness over a 12-year period to two referral centres in Marseille, in order to compare with other international data on ill returning

travellers and to inform public health surveillance and local prevention practices.

Material and Method

Study design and procedures

The GeoSentinel Surveillance Network is a global clinic based surveillance system to track travel-related illnesses within 66 clinics in 29 countries.⁴ Physicians in travel/tropical medicine electronically submit demographic, clinical and diagnostic information on patients seeking medical care for a travel-related illness having crossed an international border within the past 10 years.⁵ Every patient has at least one confirmed or probable diagnostic code, and may be assigned several. Diagnoses are based on recognition of a syndrome and/or a specific pathogen. Other data includes travel characteristics, in/outpatient status, and whether a pre-travel consultation took place.

We report all patients recorded on the GeoSentinel database in Marseille over a 12-year period. Eligible patients had been referred to two hospitals in Marseille, 'Assistance Publique—Hôpitaux de Marseille' from March 2003 to October 2015 and 'Hôpital d'Instruction des Armées Laveran' from May 2007 to October 2015.

Statistical analysis

Two time periods of interest were compared, 2003–10 and 2011–15. All summary descriptive statistics were carried out using Microsoft Access and R statistical software program. Continuous variables were described with mean and SD or median and IQR as appropriate. Categorical variables were described with numbers and percent. Unpaired *t*-tests or Wilcoxon signed rank test was used to compare continuous quantitative variables, depending on the distribution. Non-parametric Fisher tests were used to compare qualitative variables. Due to multiple tests being performed a conservative two-sided significance level of $P < 0.01$ was chosen. Time series analysis of malaria cases (the most frequent diagnosis) was performed and multiplicative decomposition of monthly data was used to examine the trend, after removing seasonality and random error.

Results

A total of 3460 patients were included during the 12-year study period with a M/F sex ratio of 1.2 and a median age of 38 years (range: 1–86 years) (Table 1). Tourism was the main reason for travel (1591/3460, 46%), followed by visiting friends or relatives (VFR) (895, 26%). The mean travel duration was 29 days and 35% (1212) of all travellers reported a pre-travel health consultation. Among 1931 travellers (56%) to Africa, 557 (16%) travelled to the Comoros archipelago alone, 1632 to sub-Saharan Africa, and 299 (9%) to North Africa. Only 634 (18%) travelled to Asia, 407 (12%) to Latin America and 136 (4%) to Europe (Figure 1). The majority of patients were seen after travel (3264, 94%) and 38% (1326) were hospitalized (Table 1). No deaths were recorded. The percentage of hospitalized patients was highest amongst travellers to sub-Saharan Africa (858/1632, 53%), with similar rates seen in the Middle East (41/75, 55%, $P = 0.8$) and North America (42/66, 64%, $P = 0.1$), but much lower rates when compared to travellers

from Asia (122/634, $P < 0.001$), Central America (46/202, $P < 0.001$) and Europe (35/136, $P < 0.001$). Higher proportions of patients visiting friends and family (573/895, 64%) were hospitalized, in comparison to tourists (409/1591, 26%, $P < 0.001$) or business travellers (163/488, 33%, $P < 0.001$). The majority of VFRs had visited sub-Saharan Africa (689/895, 77%) and did not report any pre-travel consultation (561/895, 63%).

Comparing patient characteristics between the two periods (2003–10 and 2011–15), there was a significant difference in the age of travellers, with an increase in the median age from 37 to 40. There was a significant decline in VFR as the reason for travel (599/2144, 27.94% to 296/1316, 22.49%). There was a decrease in hospitalization rates between the two periods (908/2144, 42.35% to 418/1316, 31.76%). Travel trends between the two periods differed, with increasing travel to the Caribbean (71/2144, 3.31% to 75/1316, 5.7%), the Middle East (26/2144, 1.21% to 49/1316, 3.72%), South East Asia (204/2144, 9.51% to 223/1316, 16.95%). Declining rates of travellers were seen from North America and sub-Saharan Africa (Table 1).

The most common syndromic diagnoses were febrile systemic illness (1343/3460, 39%), dermatologic (716, 21%), gastrointestinal (340, 10%) and respiratory/ear–nose–throat (331, ENT) (10%) (Figure 2A). A large number of respiratory infections (148/331, 44.7%) and acute diarrhoeal cases were undocumented (105/267, 39.3%). In comparison, the proportion of fever cases without an etiological cause was lower (209/1356, 15.6%). Unspecified skin infections and arthropod bites accounted respectively for 17.2% ($n = 125$) and 7.4% ($n = 54$) of dermatological presentations. Influenza-like illness (ILI), pneumonia, bronchitis and unspecified pharyngitis accounted respectively for 44.7% ($n = 148$), 29.6% ($n = 98$), 8.2% ($n = 27$) and 5.1% ($n = 17$) of respiratory and ENT presentations. Confirmed influenza cases included 28 influenza A, 12 H1N1 and eight influenza B. Overall, 49 patients with acute diarrhoea had confirmed²⁵ or suspected²⁴ *Giardia* infection.

The top 20 etiological diagnoses are presented in Figure 2B. Parasitic infections were largely predominant and disproportionately dominated by malaria cases followed distantly by intestinal protozoal and helminthic infections, skin parasitic infections (cutaneous larva migrans, filariasis, myiasis, leishmaniasis and scabies) and schistosomiasis. Arthropod-borne viral infections including dengue and chikungunya were also frequent. Influenza virus was the most frequently identified pathogen responsible for respiratory tract infections, followed by tuberculosis. Viral hepatitis, salmonella and cytomegalovirus infections were also among the most common diagnoses. In total, 19 rickettsial infections were identified (Figure 3).

Less frequent etiological diagnoses included bacterial infections ($n = 60$), parasitic and fungal infections ($n = 32$) and viral infections ($n = 26$) (Table 2).

The most common etiological diagnoses in patients presenting with a systemic febrile illness ($n = 1343$) were malaria (797, 59.3%), dengue (96, 7.1%), chikungunya (75, 5.5%) and cytomegalovirus (24, 1.8%). Among 797 diagnoses of malaria, *Plasmodium falciparum* accounted for 603 (76%) cases, *P. vivax* for 82 (10%), *P. ovale* for 50 (6%) and *P. malariae* for 24 (3%), while 38 were unspecified. Severe malaria affected 33 of these patients. Most malaria cases were in migrant VFRs (474/797, 59.5%) and were acquired in sub-Saharan Africa

Table 1. Overall patient demographics and travel characteristics ($n = 3460$), and comparisons between two periods (2003–10 and 2011–15)

	N (%)	2003–10, $n = 2144$	2011–15, $n = 1316$
Gender			
Male	1852 (53.53)	1158 (54.01)	694 (52.74)
Female	1598 (46.18)	977 (45.57)	621 (47.19)
Not documented	10 (0.29)	9 (0.42)	1 (0.08)
Median age in years (IQR)	38 (27–52)	39 (27–51)*	40 (28–54)*
Country of birth			
France	2324 (67.17%)	1369 (63.85)	955 (72.57)
Reason for travel			
Tourism	1591 (45.98)	958 (44.68)	633 (48.10)
Visiting friends and relatives	895 (25.87)	599 (27.94)*	296 (22.49)*
Business	488 (14.1)	286 (13.34)	202 (15.35)
Missionary/volunteer/researcher/aid work	197 (5.69)	128 (5.97)	69 (5.24)
Military	122 (3.53)	67 (3.12)	55 (4.18)
Immigration	68 (2.91)	34 (1.59)	17 (1.29)
Student	60 (1.74)	48 (2.24)*	12 (0.91)*
Medical tourism	34 (0.98)	19 (0.89)	13 (0.99)
Not documented	5 (0.14)	5 (0.23)	0
Travel duration (median, IQR)	29 (15–61)	29 (15–61)	27 (15–60)
Pre-travel encounter			
Yes	1212 (35.03)	782 (36.47)	430 (32.67)
No	1672 (48.32)	999 (46.6)	673 (51.14)
Don't know	576 (16.65)	327 (15.25)	213 (16.19)
Clinical setting			
Immigration travel only	68 (1.96)	34 (1.59)	17 (1.29)
Seen during travel	126 (3.64)	86 (4.01)	40 (3.04)
Seen after travel	3264 (94.34)	2022 (94.31)	1242 (94.38)
Not documented	2 (0.06)	2 (0.09)	0
Patient type			
Inpatient	1326 (38.32)	908 (42.35)*	418 (31.76)*
Outpatient	2124 (61.39)	1226 (57.18)*	898 (68.24)*
Not documented	10 (0.29)	10 (0.47)	0
Region of exposure			
Australia/New Zealand	6 (0.17)	6 (0.28)	0 (0)
Caribbean	146 (4.22)	71 (3.31)*	75 (5.7)*
Central America	56 (1.62)	35 (1.63)	21 (1.6)
Eastern Europe	40 (1.16)	21 (0.98)	19 (1.44)
Middle East	75 (2.17)	26 (1.21)*	49 (3.72)*
North Africa	299 (8.64)	168 (7.84)	131 (9.95)
North America	66 (1.91)	56 (2.61)*	10 (0.76)*
North East Asia	43 (1.24)	31 (1.45)	12 (0.91)
Oceania	35 (1.01)	15 (0.7)	20 (1.52)
South America	205 (5.92)	123 (5.74)	82 (6.23)
South Central Asia	164 (4.74)	102 (4.76)	62 (4.71)
South East Asia	427 (12.34)	204 (9.51)*	223 (16.95)*
Sub-Saharan Africa	1632 (47.17)	1129 (52.66)*	503 (38.22)*
Western Europe	96 (2.77)	69 (3.22)	27 (2.05)
Not ascertainable	170 (4.91)	88 (4.1)*	82 (6.23)*
Presenting syndromes			
Fever	1343 (38.82)	903 (42.12)	440 (33.43)*
Respiratory/ENT	331 (9.57)	202 (9.42)	129 (9.80)
Gastrointestinal (except acute diarrhoea)	340 (9.83)	173 (8.07)	167 (12.69)*
Acute diarrhoea	267 (7.72)	203 (9.47)	64 (4.86)*
Dermatological	716 (20.69)	447 (20.85)	269 (20.44)
Etiological conditions			
Malaria	797 (23.03)	607 (28.31)*	190 (14.44)*
Dengue	96 (2.77)	58 (2.71)	38 (2.89)
Chikungunya	75 (2.17)	29 (1.35)*	46 (3.50)*
ILI	148 (4.28)	109 (5.08)*	39 (2.96)*
Giardia	49 (1.42)	33 (1.54)	16 (1.22)

*Significant difference, at the alpha 0.01 level.

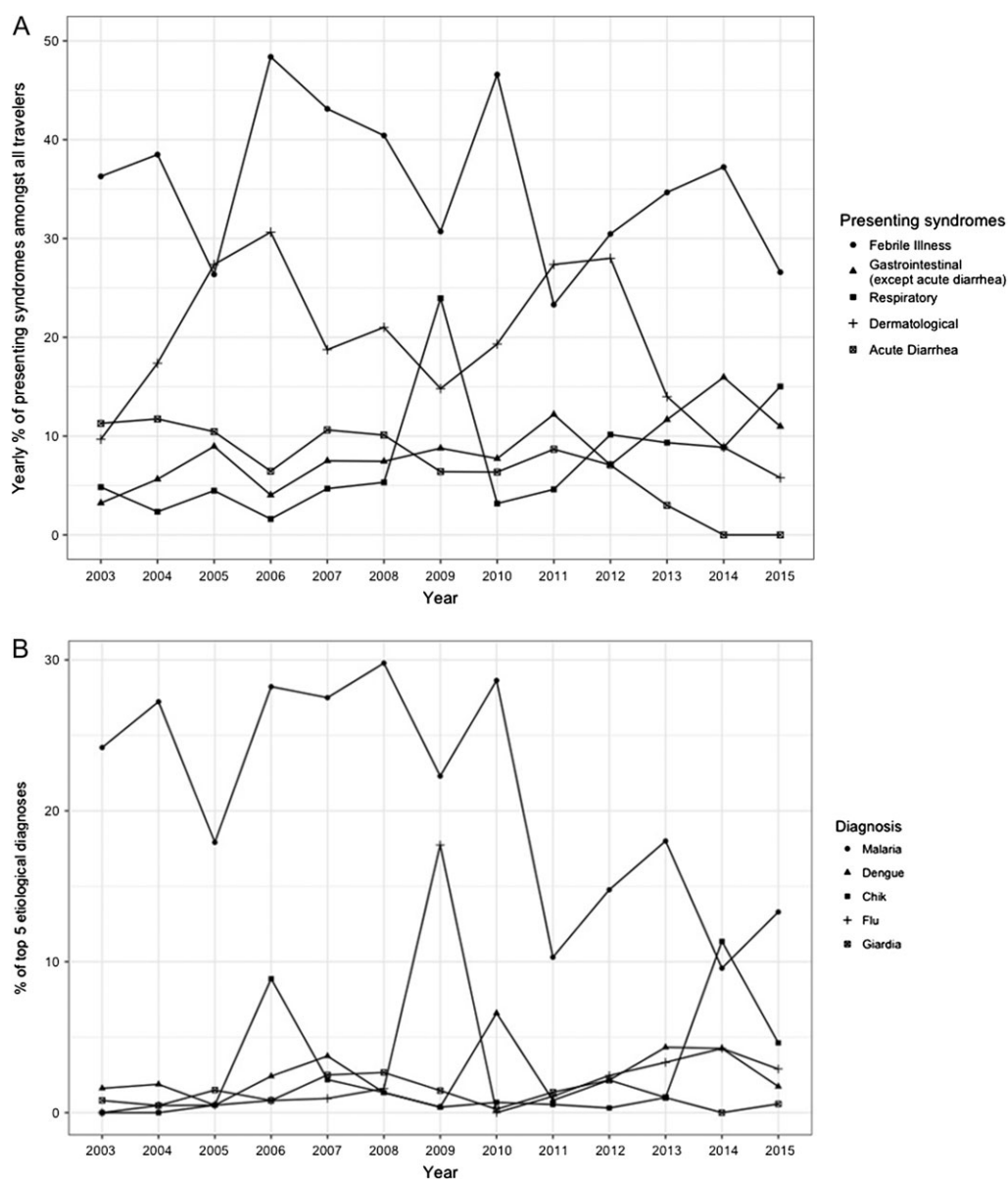


Figure 1. (A) Main rates of presenting syndromes by year 2003–2015 in travellers returning to Marseille. (B) Main rates of etiological diagnosis for Malaria, Dengue, Chikungunya, Influenza-like illness and Giardia by years 2003–15 in travellers returning to Marseille.

(729, 91.5%), notably in the Comoros archipelago (411, 51.6%) (Table 3). Figure 3 compares the overall trends after multiplicative decomposition of monthly malaria cases in the Comoros archipelago and in other exposure regions.

A total of 96 dengue cases, including two severe cases, and 75 chikungunya cases were recorded. Most cases were in French-born tourists returning from Asia, the French Caribbean islands and Africa (Table 3). Most patients with malaria, dengue and chikungunya infections did not seek pre-travel advice (409, 51.32%; 51, 53.12% and 53, 70.67%, respectively).

Regarding the presenting syndromes and diagnoses between the two periods (2003–10 and 2011–15), there was a significant increase in the number of patients with gastrointestinal symptoms, apart from acute diarrhoea, (173/2144, 8.07% to 167/1316, 12.69%) with a decrease in fever (903/2144, 42.12% to 440/1316, 33.43%), and acute diarrhoeal presentations (203/

2144, 9.47% to 64/1316, 4.86%) (Table 1). There was a significant decrease in patients with malaria (607/2144, 28.31% to 190/1316, 14.44%) and ILI (109/2144, 5.08% to 39/1316, 2.96%) between the two periods with a significant increase in the rates of chikungunya (29/2144, 1.25% to 46/1316, 3.50%).

Discussion

Many demographic characteristics reported here are comparable to global GeoSentinel surveillance studies amongst returning travellers from the developing world and in travellers returning to the USA. Similar rates in terms of sex, median age, travel duration and predominance for tourism as the reason of travel are reported.^{6,7} However, our results are characterized by the predominance of travellers returning from Africa, in comparison to Asia in a previous global GeoSentinel study.⁴ This can be

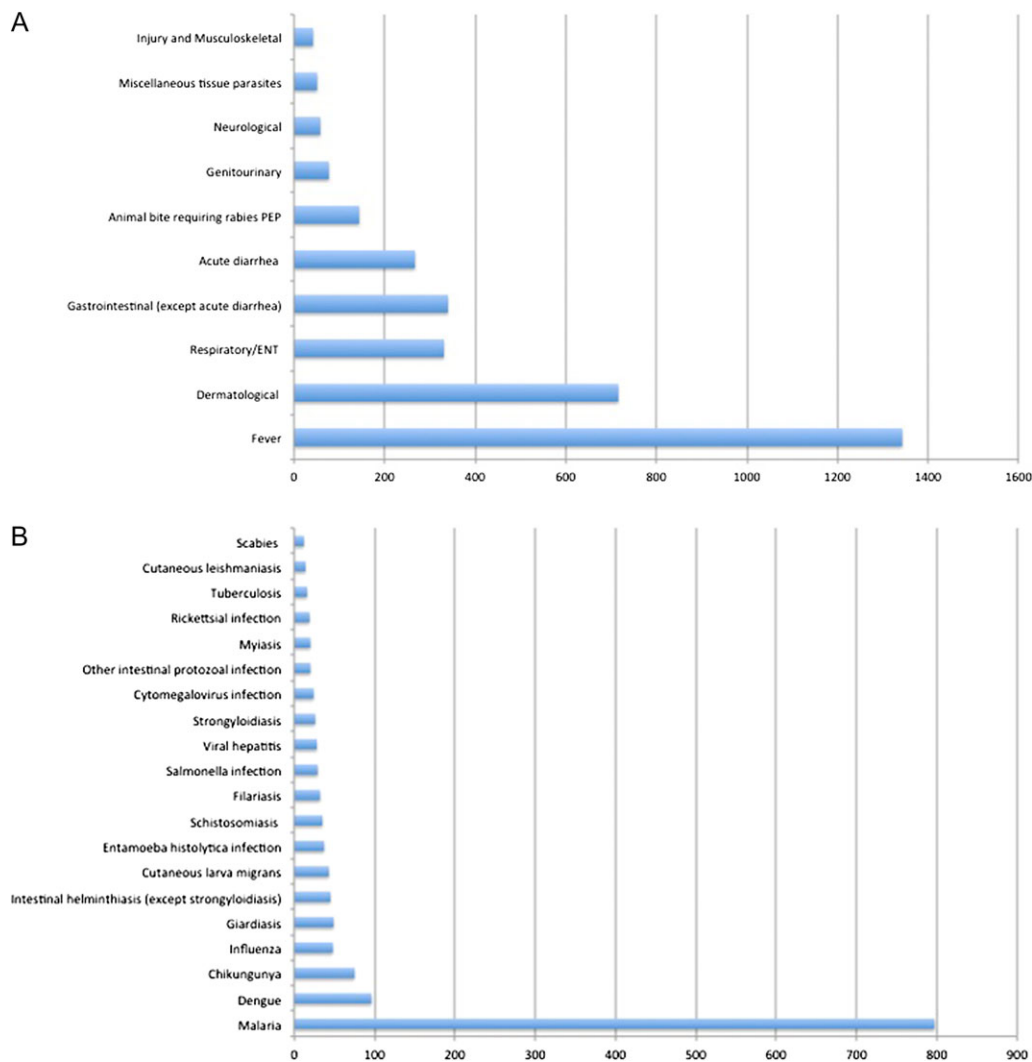


Figure 2. (A) Top 10 syndromic diagnoses ($n = 3369$ of 3794 diagnoses). (B) Top 20 diagnosis with etiologic agent identified ($n = 1465$). Further information: Dengue (uncomplicated, 94, severe 2), Influenza (28 confirmed cases of influenza A, 12 H1N1 and eight influenza B), intestinal helminthic infections (*Ascaris*, 2; Pinworm, 1; *Heterophyes*, 1; hookworm, 12; tapeworm, 4; whipworm, 2; unspecified, 23), schistosomiasis (*S. haematobium*; 9, *S. mansoni*, 15, species unknown, 11), filariasis (Bancroft, 1; loiasis, 26, other, 1; species unknown, 4), viral hepatitis (hepatitis A acute, 13; hepatitis B carrier, 1; hepatitis B acute, 2; hepatitis B chronic, 2, hepatitis C chronic, 3, hepatitis E, 7), Salmonella infections (*S. paratyphi*, 8; *S. typhi*, 8, other, 13), tuberculosis (pulmonary, 13; extrapulmonary, 3; MDR or XDR, 5), rickettsial infection (*Rickettsia orientia*, 1; tickborne spotted fever, 12; murine typhus, 2, other, 4), other intestinal protozoal infections (amoebas other than *Entamoeba histolytica*, 7; *Blastocystis*, 5; *Dientamoeba*, 2; *Isoospora*, 1; other (no further precision), 5).

explained by the local demographics in Marseille, particularly related to the migrant African population. Marseille is historically associated with Comoros migrants, with a large Comoros-originated population estimated at 50 000–70 000 in early 2000s, although more recent figures are unknown.³ This results in differences of traveller characteristics in comparison to other GeoSentinel sites, notably higher rates of VFRs, more frequent hospitalizations and low levels of pre-travel consultations. We report a disproportionately high rate of hospitalizations in patients from sub-Saharan Africa and amongst VFRs. This highlights the need to target our travel health prevention messages towards VFRs, a sub-group with known barriers to accessing pre-travel health advice.^{8,9} The African migrant population in Marseille may have specific barriers to pre-travel consultations, including financial constraints for malaria prophylaxis and lack of information. Potential strategies include primary care

physicians routinely questioning immigrant patients about future travel plans.⁸

Sentinel surveillance amongst travellers can be used to collect important epidemiological and virological information to alert international authorities to the changing epidemiology of infections, especially in countries with limited active public health surveillance.^{10,11} This is demonstrated by the decline of malaria in the Comoros. We confirm that *P. falciparum* malaria remains the most frequent cause of febrile illness in international travellers returning to Marseille, particularly from sub-Saharan Africa. This is similar to both global studies⁴ and previous local studies in Marseille from 1999 to 2003.¹² An overall decreasing trend in the number of monthly malaria cases can be noted from 2007 to 2015 (Figure 3). This is consistent with a global general decreasing trend since 2000.⁴ However, in our study, the sudden drop in 2011 is most likely due to a decrease in the

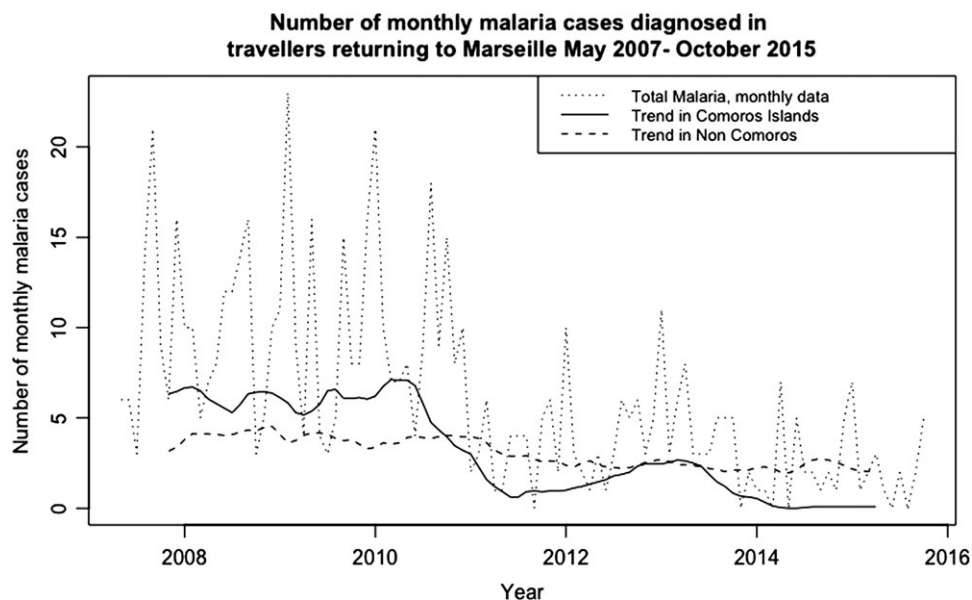


Figure 3. Number of monthly malaria cases by region of exposure. Black dotted line: monthly number of total malaria cases reported; red line: overall trend after multiplicative decomposition in Comoros; green line: overall trend after multiplicative decomposition in regions of exposure other than the Comoros.

Table 2. Less frequent infections acquired by travellers, according to main presenting complaints and type of pathogen

Presenting complaints	Bacterial infections	Parasitic and fungal infections	Viral infections
Respiratory/URTI	Legionnaire disease (4) Pertussis (2) <i>Streptococcus pharyngitis</i> (2)		
Gastrointestinal	<i>Shigella</i> (6) <i>Campylobacter</i> infection (2) <i>Helicobacter pylori</i> infection (2) <i>Clostridium difficile</i> infection (1)	Echinococcosis (5)	
Dermatological	Lyme disease (7)	Tungasis (1)	Herpes simplex infection (7) Chickenpox (4) Herpes zoster (2) Measles (2) Mumps (1) Zika virus infection (1) Acute HIV infection (4) AIDS (1)
STI	Syphilis (3) <i>Chlamydiae trachomatis</i> genital infection (2) Gonorrhoea (2)		
Neurological	Meningococcal meningitis and sepsis (2) Pneumococcal meningitis (1)	Neurocysticercosis (4) African trypanosomiasis (1)	Japanese encephalitis (1)
Wide range of symptoms	Leptospirosis (9) Q fever (8) Meliodosis (1)	<i>Toxocara</i> infection (7) Toxoplasmosis (1) Trichinellosis (1)	Epstein-Barr virus infection (3)

numbers of cases acquired in the Comoros archipelago (Figure 3); decomposition of monthly non-Comoros malaria cases demonstrated a gradual decreasing trend in comparison to the sudden drop of cases in Comoros travellers from 2011. This is in line with local malaria surveillance data demonstrating a major decline in malaria morbidity and mortality in the Union of Comoros between 2010 and 2014, which is attributed to malaria prevention and control interventions.¹³ This is an example of a local single site analysis in Marseille providing accurate sentinel surveillance information for malaria

epidemiology in Comoros. Alongside sentinel analysis for diagnosis, important information on rates of malarial drug resistance can also be collected from travellers.¹⁴

Our results highlight VFRs as disproportionately at risk for malaria, which corroborates recent French national data and from a London tropical hospital.^{15,16} Other at risk groups in our series were business travellers and military personnel, as previously observed.¹⁷⁻¹⁹

Other examples of the use of travellers as sentinels for public health surveillance includes the identification of dengue outbreaks

Table 3. Characteristics of the five main etiological diagnoses

	Malaria (N = 797)	Dengue (N = 96)	Chikungunya (N = 75)	Influenza-like illness (N = 148)	Giardia (N = 49)
Male (%)	522 (65.5)	49 (51.04)	35 (46.67)	71 (47.97)	24 (48.98)
Median age (IQR)	26 (17–36)	19 (13–29.5)	23 (12.25–31.75)	32 (1–62)	11 (5–27)
French born	254 (31.87%)	79 (82.29)	67 (89.33)	102 (68.92)	46 (93.88)
Top 3 reason for travel, N (%)	Visiting Friends and Relatives 474 (59.47) Business 109 (13.67) Military 68 (8.53)	Tourism 60 (62.5) Visiting Friends and Relatives 16 (16.67) Business 13 (13.54)	Tourism 45 (60.00) Business 15 (20) Visiting Friends and Relatives 8 (10.67)	Tourism 108 (72.97) Visiting Friends and Relatives 19 (12.84) Business 17 (11.49)	Tourism 21 (42.86) Business 11 (22.45) Student = 7 (14.29) Missionary/volunteer/research/aid work = 7 (14.29)
Top 3 regions of exposure, N (%)	Sub-Saharan Africa 729 (91.47) South America 51 (6.4) Asia 6 (0.75)	Asia 34 (35.42) Caribbean 24 (25) Sub-Saharan Africa 21 (21.88)	Caribbean 33 (44) Sub-Saharan Africa 29 (38.67) Asia 10 (13.33)	North America 36 (24.32) Western Europe 25 (16.89) Sub-Saharan Africa 22 (14.86)	Sub-Saharan Africa 16 (32.65) Asia 15 (30.61) South America 6 (12.24)
Top 5 countries of exposure, N (%)	Comoros 411 (51.57) Côte d'Ivoire 66 (8.28) French Guana 44 (5.52) Cameroon 35 (4.39) Burkina Faso 33 (4.14)	Guadeloupe 15 (15.62) Indonesia 11 (11.46) Thailand 11 (11.46) Comoros 9 (9.38) Martinique 8 (8.33)	Martinique 17 (22.67) Reunion 17 (22.67) Guadeloupe 11 (14.67) Madagascar 4 (5.33) India/Indonesia/Mauritius/Saint Martin = 3 (4)	United States 35 (23.65) Saudi Arabia 21 (14.19) Spain 14 (9.46) Comoros 11 (7.43) United Kingdom 9 (6.08)	India 10 (20.41) Madagascar 5 (10.20) Benin 3 (6.12) Burkina Faso 3 (6.12) China 3 (6.12) Indonesia 3 (6.12) Senegal 3 (6.12)
Pre-travel encounter	No 409 (51.32) Yes 305 (38.27) Don't know (9.54)	No 51 (53.12) Yes 29 (30.21) Don't know 16 (16.67)	No 53 (70.67) Yes 16 (21.33) Don't know 6 (8)	No 59 (39.86) Yes 57 (38.51) Don't know 32 (21.62)	No 12 (24.49) Yes 23 (46.94) Don't know (26.53)

in Luanda and Dar es Salaam¹¹ and for the Indian Ocean and West Africa.^{20,21} Due to a large number of our patients returning from Africa, travellers returning to Marseille may be useful sentinels for future arboviral African outbreaks. Rapid testing for dengue is essential in patients presenting with a fever from Africa. Similarly, the geographical profile and number of travellers presenting with chikungunya in Marseille (Table 3 and Figure 1B) is a reflection of increasing global expansion, providing further examples of vital sentinel surveillance in travellers.^{22,23} As our survey ended in October 2015, we only report one case of Zika virus; however, this is now an important differential amongst febrile travellers.²⁴ Marseille may be a vital sentinel site to document possible Zika epidemics on the African continent.²⁵

Public health surveillance of these arboviral infections are of particular interest in the Mediterranean region, due to the local emergence of arthropod-borne diseases given the presence of the *Aedes albopictus* vector in the Mediterranean region and risk of autochthonous transmission, particularly around hospitals.²⁶ Arthropod-borne diseases accounted for 68% of the top 20 diagnoses. Dengue was the second most common diagnosis during our study period, with 35% of patients from Asia, consistently with global data.^{4,7} Furthermore, we report a significant increase in chikungunya cases between the two periods. This high rate of imported arboviral diseases is likely to increase, given the trend of increasing numbers of unwell travellers returning from the Caribbean and South East Asia. This indicates the need for improved prevention strategies, both before travel, by reinforcing pre-travel advice toward prevention of insect bites and malaria chemoprophylaxis, and after, with improved epidemiological surveillance and healthcare professional training. Pre- and post-travel advice should also include advice on reducing respiratory transmission (such as local emergence of multi-resistant tuberculosis or Middle East Respiratory Syndrome (MERS) coronavirus), and improved hygiene practices to reduce nosocomial transmission of multi-resistant bacteria. Furthermore, improved diagnostic tools after travel, such as multiplex PCR, may help increase the number of etiological diagnoses amongst syndromic diarrhoeal and respiratory illnesses.

Another specificity of Marseille are the large numbers of Muslim Hajj pilgrims each year returning from Mecca.²⁷ We demonstrate the increasing trend of travellers returning from the Middle East, demonstrating the importance of Marseille as a sentinel surveillance site to the Middle East. Over the last few years, the vast majority of travellers with respiratory symptoms after returning from the Middle East consulted because of a fear of contracting MERS.²⁸ Due to a protocolized management procedure, patients returning from the Middle East ($n = 75$) were disproportionately hospitalized (41, 55%), resulting in a hospitalization bias.

Finally, the characteristics of sentinel surveillance in Marseille are also related to the specifics of the medical community. We house the national reference centre for rickettsioses, and so provide an opportunity to document high risk geographic areas for these emerging, underestimated infections.²⁹ Similarly, the centre for rabies vaccination is situated in Marseille, providing sentinel surveillance on patients seeking rabies post-exposure prophylaxis. We highlight the potential for rabid animal exposure in travellers, notably in Asia and the need to reinforce pre-exposure prophylaxis.³⁰

Previous studies local to Marseille have used a prospective cohort design, providing a denominator, amongst Hajj pilgrims³¹ and travellers to Senegal.³² This approach allows an estimation of risk of travel-associated diseases, including mild self-limiting conditions but is biased by traveller selection. Here, we use a different approach that describes the spectrum of diseases in patients returning from any foreign destinations and being referred to a specialized structure. Limitations include an important recruitment bias, with an underestimate of self-limiting conditions, a lack of denominator and disease prevalence estimates. However, it captures more severe and uncommon travel-related illnesses and provides vital sentinel information on the changing epidemiology of infectious diseases across the globe, most notably for dengue and chikungunya, as well as identifying less common infectious disease outbreaks in the tropics where laboratory facilities may be limited.³³

In conclusion, we demonstrate the need to understand the local population and corresponding travel risks, particular amongst those visiting friends and relatives who have higher hospitalization rates post-travel. Sentinel surveillance amongst travellers in Marseille provides essential information to guide both local practices, especially pre-travel consultation needs and addressing autochthonous vector-borne risks, and informing institutional partners worldwide regarding the spread of arboviral epidemics and other arthropod borne infections.

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Author contributions

K.G. analysed the data and wrote the article. H.S., F.S., P.B., P.P. collected data, and edited the article. P.G. collected and analysed the data and wrote the article.

References

1. Direction Générale des Entreprises. *Etudes économiques*. Chiffres clés du tourisme [Internet]. 2016 [cited 2017 Oct 6]. http://www.entreprises.gouv.fr/files/files/directions_services/etudes-et-statistiques/stats-tourisme/chiffres-cles/2016-Chiffres-cles-tourisme-FR.pdf
2. Bertonecello B, Bredeloup S. Le Marseille des marins africains. *Rev Eur Migr Int* 1999; 15(3):177–97.
3. Parola P, Gazin P, Pradines B *et al*. Marseilles: a surveillance site for malaria from the Comoros Islands. *J Travel Med* 2004; 11(3): 184–6.

4. Leder K, Torresi J, Brownstein JS *et al.* Travel-associated illness trends and clusters, 2000–2010. *Emerg Infect Dis* 2013; **19**(7):1049–73.
5. Harvey K, Esposito DH, Han P *et al.* Surveillance for travel-related disease—GeoSentinel Surveillance System, United States, 1997–2011. *Morb Mortal Wkly Rep Surveill Summ Wash DC* 2002 2013; **62**: 1–23.
6. Hagmann SHF, Han PV, Stauffer WM *et al.* Travel-associated disease among US residents visiting US GeoSentinel clinics after return from international travel. *Fam Pract* 2014; **31**():678–87.
7. Freedman DO, Weld LH, Kozarsky PE *et al.* Spectrum of disease and relation to place of exposure among ill returned travelers. *N Engl J Med* 2006; **354**(2):119–30.
8. Leder K, Lau S, Leggat P. Innovative community-based initiatives to engage VFR travelers. *Travel Med Infect Dis* 2011; **9**(5):258–61.
9. Seale H, Kaur R, Mahimbo A *et al.* Improving the uptake of pre-travel health advice amongst migrant Australians: exploring the attitudes of primary care providers and migrant community groups. *BMC Infect Dis [Internet]* 2016 [cited 2017 Jun 20]; **16**(1). <http://bmcinfectdis.biomedcentral.com/articles/10.1186/s12879-016-1479-1>.
10. Gautret P, Botelho-Nevers E, Charrel RN, Parola P. Dengue virus infections in travellers returning from Benin to France, July–August 2010. *Euro Surveill Bull Eur Sur Mal Transm Eur Commun Dis Bull* 2010; **15**(36):pii: 19657:1–2.
11. Neumayr A, Munoz J, Schunk M *et al.* Sentinel surveillance of imported dengue via travellers to Europe 2012 to 2014: TropNet data from the DengueTools Research Initiative. *Eurosurveillance [Internet]* 2017 Jan 5 [cited 2017 Jun 20]; **22**(1). <http://www.eurosurveillance.org/ViewArticle.aspx?ArticleId=22679>.
12. Parola P, Soula G, Gazin P *et al.* Fever in travelers returning from tropical areas: prospective observational study of 613 cases hospitalised in Marseilles, France, 1999–2003. *Travel Med Infect Dis* 2006; **4**(2):61–70.
13. Kassim SA, James PB, Alolga RN *et al.* Major decline in malaria morbidity and mortality in the Union of Comoros between 2010 and 2014: the effect of a combination of prevention and control measures. *South Afr Med J Suid-Afr Tydskr Vir Geneeskde* 2016; **106**(7):709–14.
14. Gharbi M, Flegg JA, Hubert V *et al.* Longitudinal study assessing the return of chloroquine susceptibility of *Plasmodium falciparum* in isolates from travellers returning from West and Central Africa, 2000–2011. *Malar J* 2013; **12**(1):35.
15. Centre National de Référence du Paludisme. *Rapport d'activité 2015 [Internet]*. Centre National de Référence du Paludisme; 2015. http://cnrpaludisme-france.org/docs/rapport_activites_cnr_paludisme_2014.pdf
16. Marks M, Armstrong M, Whitty CJM, Doherty JF. Geographical and temporal trends in imported infections from the tropics requiring inpatient care at the Hospital for Tropical Diseases, London—a 15 year study. *Trans R Soc Trop Med Hyg* 2016; **110**(8):456–63.
17. Zhou S, Li Z, Cotter C *et al.* Trends of imported malaria in China 2010–2014: analysis of surveillance data. *Malar J [Internet]* 2016 [cited 2017 Jun 20]; **15**(1). <http://www.malariajournal.com/content/15/1/39>.
18. Newman RD, Parise ME, Barber AM, Steketee RW. Malaria-related deaths among U.S. travelers, 1963–2001. *Ann Intern Med* 2004; **141**(7):547–55.
19. Rapp C, Aoun O, Ficko C *et al.* Infectious diseases related aero-medical evacuation of French soldiers in a level 4 military treatment facility: a ten year retrospective analysis. *Travel Med Infect Dis* 2014; **12**(4):355–9.
20. Gautret P, Simon F, Hervius Askling H *et al.* Dengue type 3 virus infections in European travellers returning from the Comoros and Zanzibar, February–April 2010. *Euro Surveill Bull Eur Sur Mal Transm Eur Commun Dis Bull* 2010; **15**(15):19541.
21. Ninove L, Parola P, Baronti C *et al.* Dengue virus type 3 infection in traveler returning from West Africa. *Emerg Infect Dis* 2009; **15**(11): 1871–2.
22. Wahid B, Ali A, Rafique S, Idrees M. Global expansion of chikungunya virus: mapping the 64-year history. *Int J Infect Dis*. 2017; **58**: 69–76.
23. Savini H, Gautret P, Gaudart J *et al.* Travel-associated diseases, Indian Ocean Islands, 1997–2010. *Emerg Infect Dis* 2013; **19**(8): 1297–301.
24. Hamer DH, Barbre KA, Chen LH *et al.* Travel-associated zika virus disease acquired in the Americas through February 2016: a GeoSentinel analysis. *Ann Intern Med* 2017; **166**(2):99–108.
25. Nutt C, Adams P. Zika in Africa: the invisible epidemic? *Lancet* 2017; **389**(10079):1595–6.
26. Cotteaux-Lautard C, Berenger J-M, Fusca F *et al.* A new challenge for hospitals in Southeast France: monitoring local populations of *Aedes albopictus* to prevent nosocomial transmission of dengue or chikungunya. *J Am Mosq Control Assoc* 2013; **29**(1):81–3.
27. Griffiths K, Charrel R, Lagier J-C *et al.* Infections in symptomatic travelers returning from the Arabian peninsula to France: a retrospective cross-sectional study. *Travel Med Infect Dis* 2016; **14**(4): 414–6.
28. Gautret P, Benkouiten S, Griffiths K, Sridhar S. The inevitable Hajj cough: surveillance data in French pilgrims, 2012–2014. *Travel Med Infect Dis* 2015; **13**(6):485–9.
29. Delord M, Socolovschi C, Parola P. Rickettsioses and Q fever in travelers (2004–2013). *Travel Med Infect Dis* 2014; **12**(5):443–58.
30. Gautret P, Harvey K, Pandey P *et al.* Animal-associated exposure to rabies virus among travelers, 1997–2012. *Emerg Infect Dis* 2015; **21**(4):569–77.
31. Gautret P, Soula G, Delmont J *et al.* Common health hazards in French pilgrims during the Hajj of 2007: a prospective cohort study. *J Travel Med* 2009; **16**(6):377–81.
32. Dia A, Gautret P, Adheossi E *et al.* Illness in French travelers to Senegal: prospective cohort follow-up and sentinel surveillance data. *J Travel Med* 2010; **17**(5):296–302.
33. Aubry C, Gautret P, Nougairede A *et al.* Outbreak of acute haemorrhagic conjunctivitis in Indian Ocean Islands: identification of Coxsackievirus A24 in a returned traveller. *Euro Surveill Bull Eur Sur Mal Transm Eur Commun Dis Bull*. 2012 May 31; **17**(22):1–3.