

Screening vs empirical treatment of migrants for parasitic infections

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Plan

- Key issues
- Examples of data
- Proposal

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Why screen?

- Treat patient now
- Prevent late disease
- Public health
 - Direct transmission to others
 - Food handlers
 - Protect blood supply

Screening issues

- Identify migrant groups
- When?
 - Pre departure
 - On arrival
- Framework for reception and screening
- Number and types of test
- Needs geographic knowledge
- Retention and treatment
- Cost

Treatment issues

- Mass drug treatment more effective
 - Eg albendazole for helminths
 - Or ivermectin?
- Praziquantel for schistosomiasis
- Primaquine or other for malaria
- etc

Balance

- Cost and complexity of screening

Vs

- Cost, effectiveness and side effects of treatment

- NB special groups eg

- Children

- Pregnancy

- HIV

Muennig et al NEJM 1998

- Empirical treatment

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Infectious Disease Screening for Refugees Resettled in the United States

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Refugees resettling in the United States carry a significant burden of infectious diseases as a result of exposures in their countries of origin and the circumstances of their migration. Overseas screening is required before entry, but it incompletely assesses infectious diseases in refugees. Domestic health assessment has the potential to provide more comprehensive assessment for infectious diseases. Screening protocols ideally should test for tuberculosis, hepatitis B, and intestinal and other parasites and should include mechanisms for providing or updating immunizations. Testing for other infectious diseases, including malaria, hepatitis C, human immunodeficiency virus, and sexually transmitted diseases, can be performed on the basis of clinical signs and symptoms. This article reviews the current status of overseas and domestic health screening for refugees, infectious disease burdens, and future goals for health assessment of refugees and other immigrants.

CID 2004

Table 1. Screening tests for infectious diseases recommended for all refugees.

| Test, finding | Associated disease/condition |
|-------------------------------------------------------------|------------------------------------------------------------------------------|
| Tuberculin skin test: positive skin test result | TB; nontuberculous mycobacteria |
| Hepatitis B screening | |
| HBsAb | Immune |
| HBsAg | Carrier state, current, or chronic infection |
| HBcAb | Current or past infection |
| Complete blood cell count | |
| Low WBC count | HIV infection |
| Low hemoglobin level or hematocrit | Malaria |
| Lymphopenia | HIV infection |
| Eosinophilia | Parasitic diseases (e.g., schistosomiasis, filariasis, and strongyloidiasis) |
| Thrombocytopenia | Malaria; HIV infection |
| Urinalysis | |
| Hematuria | Schistosomiasis |
| Pyuria | Urinary tract infection; renal TB |
| Microscopic evaluation of stool specimens: ova or parasites | See table 2 |

NOTE. HBcAb, antibody to hepatitis B core antigen; HBsAb, antibody to hepatitis B surface antigen; HBsAg, hepatitis B surface antigen; TB, tuberculosis.

Table 2. Common parasites in refugees.

Pathogens

Ascaris lumbricoides (roundworm)

Entamoeba histolytica

Giardia lamblia

Hookworm

Hymenolepis nana (dwarf tapeworm)

Schistosoma species

Schistosoma hematobium

Schistosoma mansoni

Schistosoma japonicum

Trichuris trichiura (whipworm)

Possible pathogens in symptomatic individuals

Blastocystis hominis

Dientamoeba fragilis

Nonpathogens

Blastocystis hominis (see above)

Chilomastix mesnili

Endolimax nana

Entamoeba coli

Entamoeba hartmanni

Iodamoeba butschlii

High Prevalence and Presumptive Treatment of Schistosomiasis and Strongyloidiasis among African Refugees

Drew L. Posey,^{1,3,a} Brian G. Blackburn,^{2,3,a,b} Michelle Weinberg,¹ Elaine W. Flagg,¹ Luis Ortega,¹ Marianna Wilson,² W. Evan Secor,² Kolby Sanders-Lewis,² Kimberly Won,² and James H. Maguire²

¹Division of Global Migration and Quarantine, National Center for Preparedness, Detection, and Control of Infectious Diseases, ²Division of Parasitic Diseases, National Center for Zoonotic, Vector-borne, and Enteric Diseases, Coordinating Center for Infectious Diseases, and ³Epidemic Intelligence Service Program, Office of Workforce and Career Development, Centers for Disease Control and Prevention, Atlanta, Georgia

CID 2007

Spectrum of Illness in International Migrants Seen at GeoSentinel Clinics in 1997–2009, Part 1: US-Bound Migrants Evaluated by Comprehensive Protocol-Based Health Assessment

Elizabeth D. Barnett,¹ Leisa H. Weld,² Anne E. McCarthy,³ Heidi So,⁴ Patricia F. Walker,⁵ William Stauffer,^{5,6} and Martin Cetron,⁶ for the GeoSentinel Surveillance Network

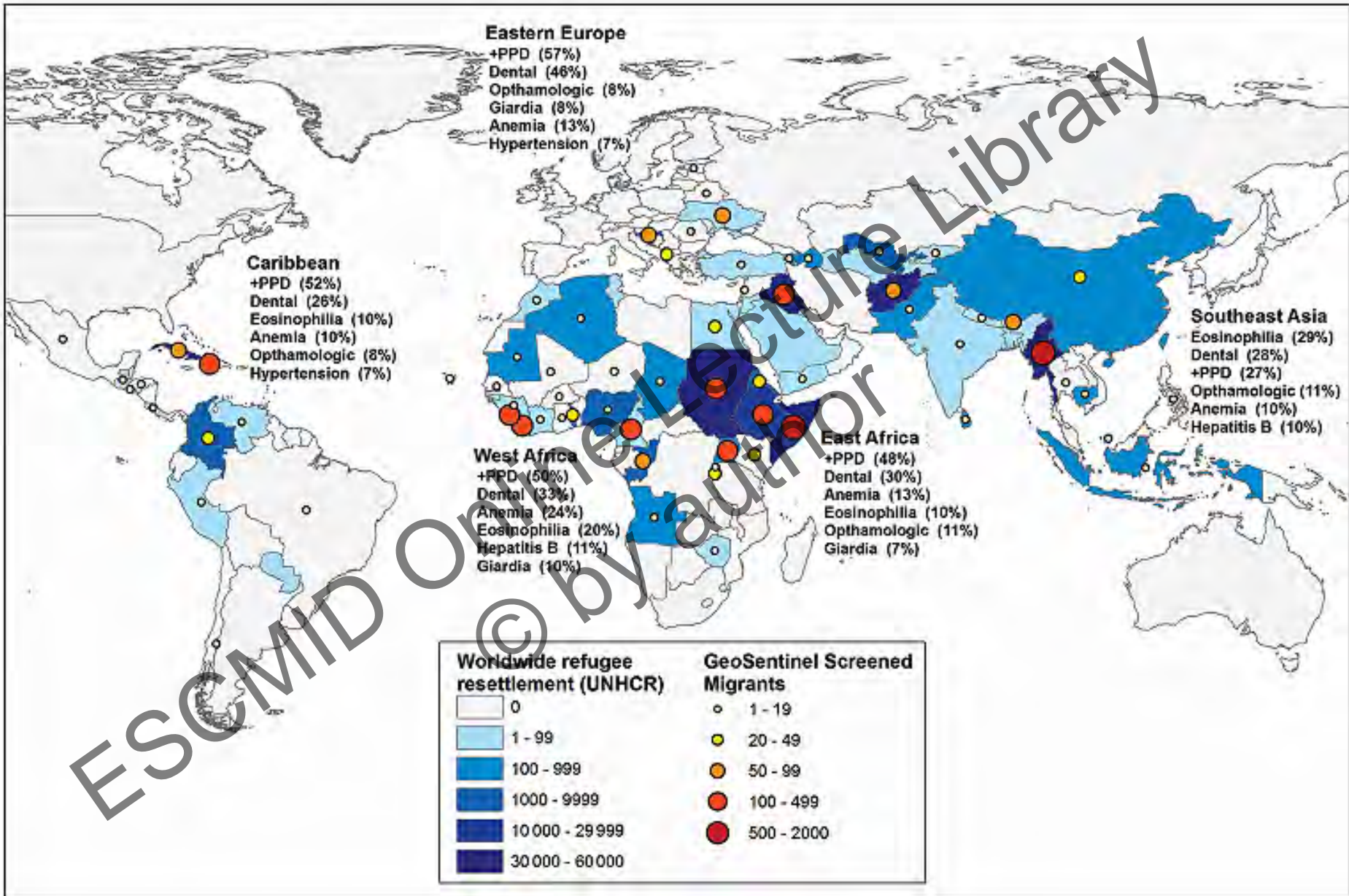
CID 2013

Table 3. Frequent Diagnoses in Migrants Evaluated by Protocol-Based Health Assessment

| Most Frequent Diagnoses | All Regions (N = 7792) | % |
|--------------------------------|------------------------|----|
| +PPD (latent tuberculosis) | 3367 | 43 |
| Dental | 2485 | 32 |
| Eosinophilia | 1191 | 15 |
| Anemia | 861 | 11 |
| Ophthalmologic | 768 | 10 |
| No health condition identified | 707 | 9 |
| Giardia | 545 | 7 |
| Hepatitis B | 497 | 6 |
| Hypertension | 379 | 5 |
| Dermatologic | 396 | 5 |
| Hematuria | 284 | 4 |

Nonpathogenic protozoa were found in 1055 (14%) of screened migrants.

Abbreviation: PPD, purified protein derivative.



Spectrum of Illness in International Migrants Seen at GeoSentinel Clinics in 1997–2009, Part 2: Migrants Resettled Internationally and Evaluated for Specific Health Concerns

Anne E. McCarthy,¹ Leisa H. Weld,³ Elizabeth D. Barnett,⁵ Heidi So,² Christina Coyle,⁶ Christina Greenaway,⁴ William Stauffer,^{8,9} Karin Leder,¹⁰ Rogelio Lopez-Velez,¹¹ Phillippe Gautret,¹² Francesco Castelli,¹³ Nancy Jenks,⁷ Patricia F. Walker,⁸ Louis Loutan,^{14,15} and Martin Cetron⁹; for the GeoSentinel Surveillance Network^a

CID 2013

Table 1. Ten Most Frequent Infectious Disease Diagnoses in Referred Migrants, Children and Adults

| Diagnosis | Frequency | Percentage |
|-------------------------------------|-----------|------------|
| Children (age ≤ 18 y; n = 854) | | |
| Malaria | 170 | 20.0 |
| Latent tuberculosis | 92 | 10.8 |
| No health condition identified | 82 | 10.0 |
| Schistosomiasis | 71 | 8.3 |
| Giardiasis | 67 | 7.8 |
| Active tuberculosis | 65 | 7.6 |
| Hepatitis B, acute and chronic | 41 | 4.8 |
| Strongyloidiasis | 40 | 4.7 |
| Eosinophilia | 25 | 2.9 |
| Intestinal ascaris | 19 | 2.2 |
| Adults (age ≥ 19 y, n = 6751) | | |
| Latent tuberculosis | 1619 | 24.0 |
| Hepatitis B, acute and chronic | 864 | 12.8 |
| Active tuberculosis | 723 | 10.7 |
| Human immunodeficiency virus/AIDS | 510 | 7.6 |
| Schistosomiasis | 370 | 5.5 |
| Hepatitis C | 346 | 5.1 |
| Strongyloidiasis | 344 | 5.1 |
| No health condition identified | 326 | 4.8 |
| Malaria | 321 | 4.8 |
| Eosinophilia | 182 | 2.7 |

NEJM 2012

Albendazole Therapy and Enteric Parasites in United States–Bound Refugees

Stephen J. Swanson, M.D., Christina R. Phares, Ph.D., Blain Mamo, M.P.H.,
Kirk E. Smith, D.V.M., Ph.D., Martin S. Cetron, M.D.,
and William M. Stauffer, M.D.

CONCLUSIONS

Presumptive albendazole therapy administered overseas before departure for the United States was associated with a decrease in the prevalence of intestinal nematodes among newly arrived African and Southeast Asian refugees.

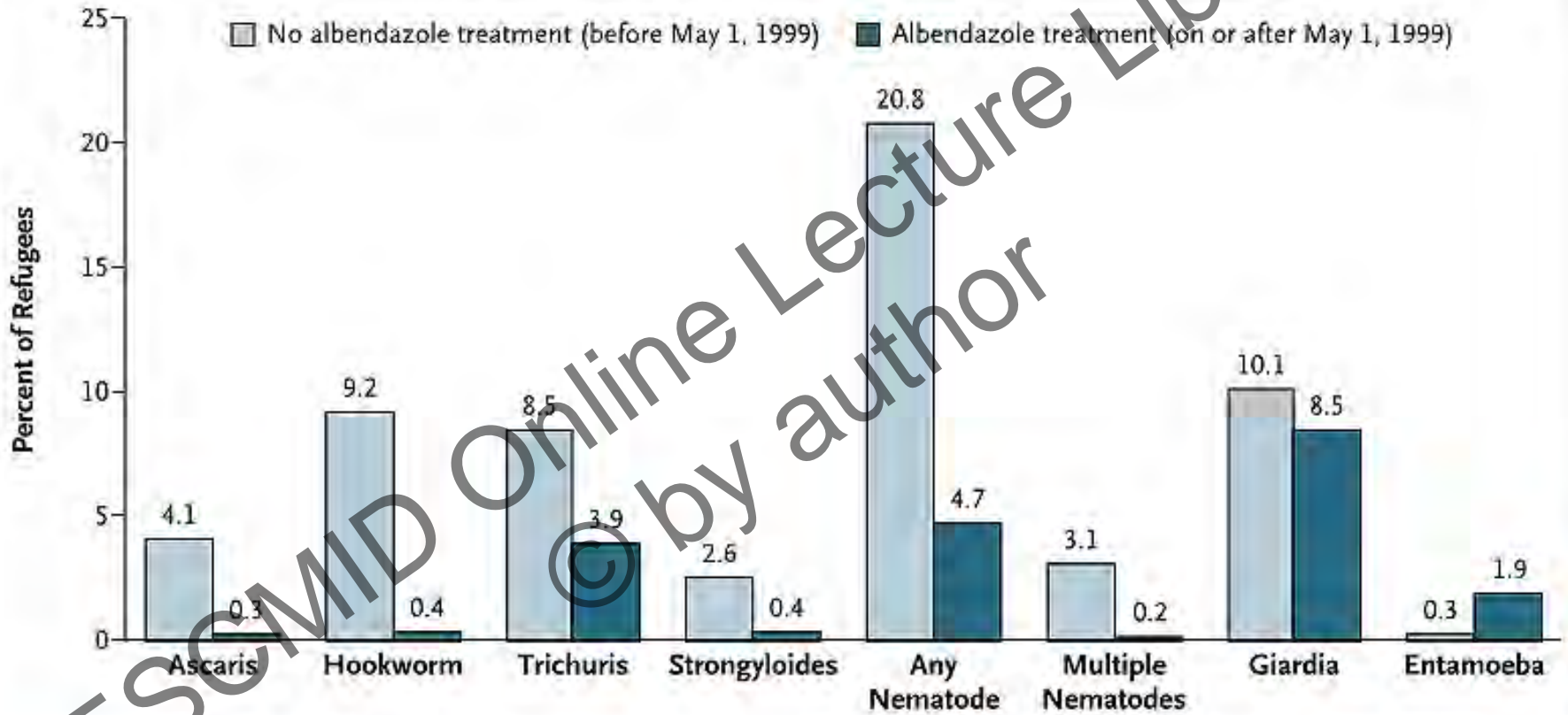


Figure 1. Prevalence of Intestinal Parasites among 26,956 Refugees Arriving in Minnesota, According to Status with Respect to Albendazole Treatment before Departure for the United States.

Proposal

- ESGITM and partners
- Literature review on European migrant parasitic infection profiles
- Identify sites with screening programmes
- Prospective & retrospective data modelling for effectiveness and cost
- Design and test appropriate screening and treatment strategies

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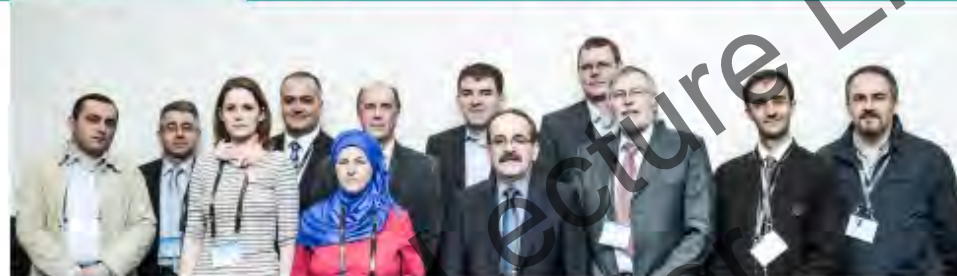
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ESCMID Study Group for Infections in Travellers and Migrants - ESGITM

News & Activities



ESGITM @ ECCMID 2014 in Barcelona, ES

- Saturday, 10 May 2014**
11.00 - 13.00 (Hall D)
Educational Workshop 13: Basics of infections in travellers.
- Sunday, 11 May 2014**
7.45 - 8.45 (Hall H)
ESGITM Meet-the-Expert Session: Controversies in treatment of malaria and Crimean-Congo haemorrhagic fever.
18.15 - 19.15 (Room 125)
ESGITM Business Meeting: All interested persons are cordially invited to join the meeting and plan future Study Group activities.

For more details, please check the ECCMID programme.

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