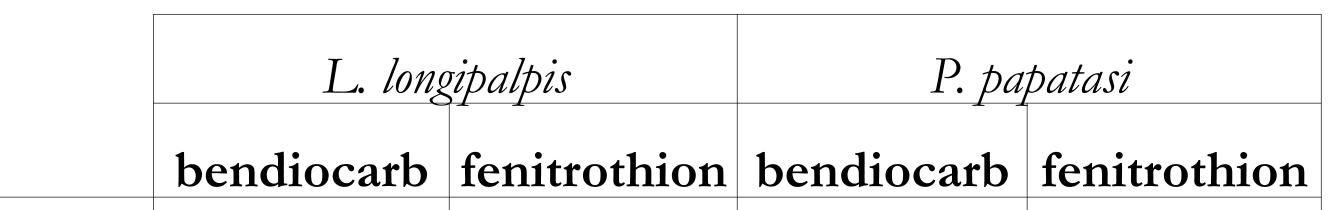


Introduction

Principal Findings

Insecticide resistance to synthetic chemical insecticides is becoming a worldwide (Fig. 3) concern in phlebotomine sand flies (Diptera: Psychodidae), the vectors of Figure 1. mucocutaneous *Leishmaniasis* infection (left), Leishmania parasites (Fig. 1 & 2). micrograph of Leishmania promastigotes parisites (right). The CDC bottle bioassay assesses resistance by testing populations against verified diagnostic doses and diagnostic times for an insecticide, but the assay has been Sandfly takes a blood meal Promastigotes are phagocytized by used limitedly with sand flies. B Divide in midgut and migrate to proboscis The objective of this study 1 was to determine diagnostic doses and diagnostic





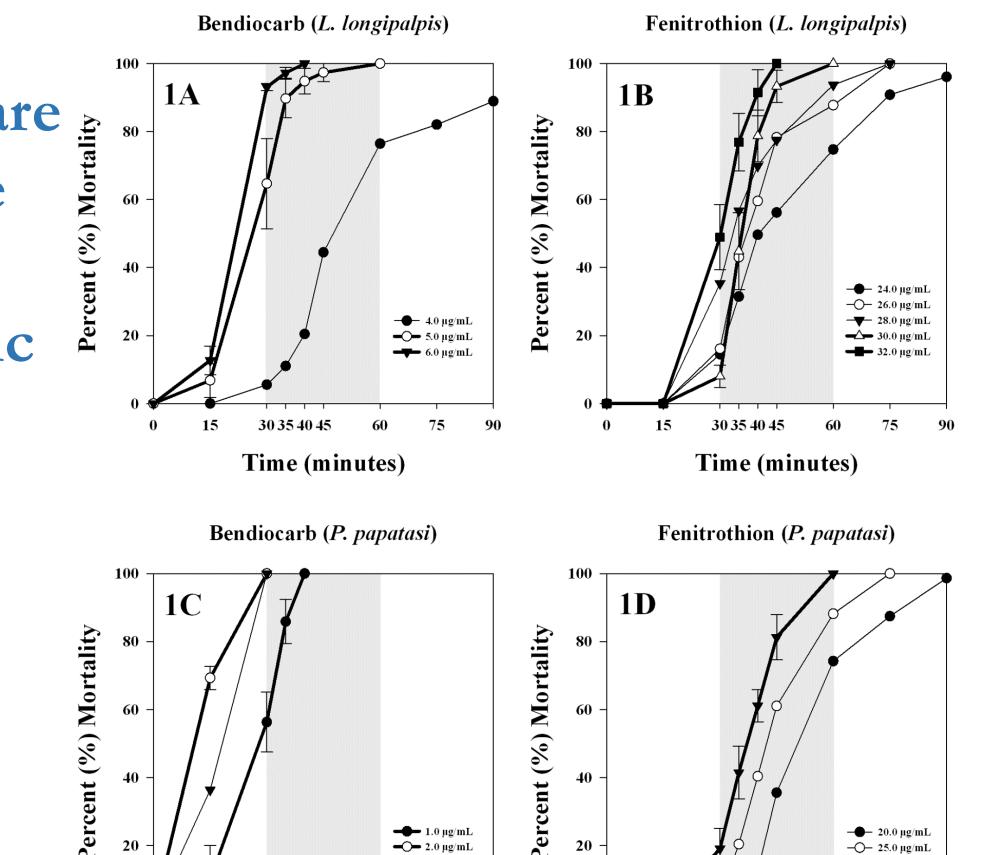
times for laboratory Lutzomyia longipalpis (Lutz and Nieva) and Phlebotncluding macrophages) of omus papatasi (Scopoli)

arious tissues 6 Ingestion of parasitized cell TDC to ten insectcides, include-= Infective Stage a = Diagnostic Stage http://www.dpd.cdc.gov/dpdx ing pyrethroids, orange-Figure 2. Leishmania life cycle phosphates, carbomates, and DDT, that are used worldwide to control sandflies (Fig. 1).

Diagnostic dose (µg/mL)	6	32	1	30
Diagnostic time (min)	40	45	40	60

Table 1. Diagnostic doses and times for bendiocarb (carbomate) and fenitroghion (organophosphate) to *L. longipalpis* and P. papatasi sand flies.

Lutzomyi longipalpis and Phlebotomous papatasi sand flies are both highly susceptible to the carbamates, such as benciocarb, as their diagnostic doses are under 7.0 µg / ml (Table 1, Fig.5). Both species are very susceptible to DDT during the exposure assay as their diagnostic doses are 7.5 µg / ml, yet their diagnostic doses for the 24-h recovery period are 650.0 μ g / ml for L. longipalpis and 470.0 µg / ml for P. papatasi.



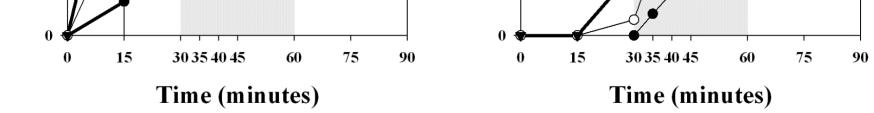


Figure 5. Diagnostic doses and times for bendiocarb (carbonate) and Fenitrothion (organophosphate) to L. *longipalpis* and P. *papatasi* sand flies.

Figure 3. Countries highlighted in red have documented cases of sand fly

tolerance or resistance to insecticides.

Conclusions/Significance

Methodology

Bioassays were conducted in 1000-ml glass bottles each containing 10-25 sand flies from laboratory colonies of L. Longipalpis or P. Papatasi (Fig. 4). Four pyrethroids, three organophosphates, two carbamates, and one organochlorine were evaluated. A range of concentrations were tested for each insecticide, and four replicates were conducted for each concentration. Diagnostic doses were determined only during the exposure bioassay for the organophosphates and carbamates. For the pyrethroids and DDT, diagnostic doses were determined for both the exposure bioassay and after a 24-hour recovery period.

Diagnostic doses and diagnostic times can now be incorporated into vector management programs that use the CDC bottle bioassay to assess insecticide resistance in wild populations of L. longipalpis



Figure 6. Female Phlebotomus papatasi taking a blood meal. and P. papatasi (Fig.6). These findings provide initial starting points for determining diagnostic doses and diagnostic times for other sand fly vector species and wild populations using the CDC bottle bioassay.

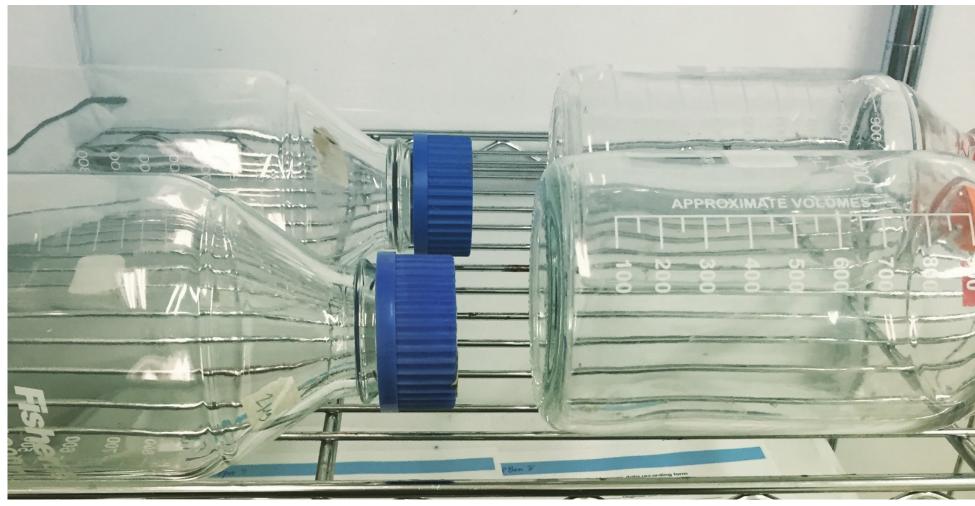


Figure 4. CDC Bottle Bioassay procedure

Amastigotes transform

7 promastigote stage in midgr



We thank Laine Anderson, Conor Reese, Michael Higham, Parker Johnson, Michael Preece, and Tess Jolley (Utah State University) for their laboratory assistance with maintaining the sand fly colonies and diagnostic dose determination assistance.