TABLE

Results for the major hits obtained after BLAST analysis of the 234 bp-hsp70C polymerase chain reaction and sequencing

Sample	Description	Max score	Total score	Query cover	E value	Max identity	Accession number
2 - Evandromyia saulensis	Leishmania braziliensis MHOM/BR/75/M2904 putative heat-shock protein hsp70 (LBRM_28_2970) mRNA, partial cds	261	261	78%	2,00E-66	96%	gi 389601962 XM_001566273.2
6 - Trichophoromyia auraensis	L. braziliensis MHOM/BR/75/M2904 putative heat-shock protein hsp70 (LBRM_28_2970) mRNA, partial cds	217	217	97%	2,00E-53	97%	gi 389601962 XM_001566273.2
3 - <i>Pressatia</i> sp	L. braziliensis MHOM/BR/75/M2904 putative heat-shock protein hsp70 (LBRM_28_2970) mRNA, partial cds	250	250	100%	3,00E-63	95%	gi 389601962 XM_001566273.2
11 - Th. auraensis	L. braziliensis MHOM/BR/75/M2904 putative heat-shock protein hsp70 (LBRM_28_2970) mRNA, partial cds	289	289	100%	6,00E-75	100%	gi 389601962 XM_001566273.2
5 - Th. auraensis	L. braziliensis strain MHOM/BR/2002/NMT-LTCP 14440-P clone B heat shock protein 70 (hsp70) gene, partial cds	379	379	100%	5,00E- 102	99%	gi 316891027 GU368187.1

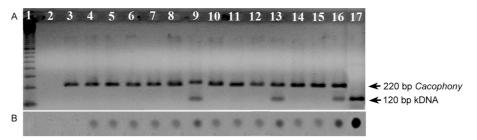


Fig. 1: natural infection by *Leishmania* spp in sandflies. (A) multiplex polymerase chain reaction (PCR) directed to the 220 bp fragment of the cacophony gene (sandflies) and the 120 bp of kDNA conserved minicircle sequence (*Leishmania* spp.) was performed individually in 173 sandfly DNA extracts. Amplified products were visualised on 2% agarose gel stained with ethidium bromide. 1: molecular weight 100 bp DNA ladder (Invitrogen); 2: negative control for PCR reaction (no DNA added); 3: negative control for DNA extraction (DNA extracted from male sandflies); 4-15: single female sandflies subjected to molecular diagnosis that revealed positivity in two samples (lines 9 and 13); 16: positive control (DNA extracted from laboratory-bred *Lutzomyia longipalpis* females); 17: positive control (DNA from cultivated *L. (V.) braziliensis*) [10 ng]. (B) dot blot hybridisation of PCR-amplified products with a specific byotinylated probe for *Leishmania* subgenus *Viannia*. 4-5: *Evandromyia saulensis* - Area of forest (area I); 6: *Pressatia* sp. - Area of forest (area II); 7-15: *Trichophoromyia auraensis* - Area of forest (area II); 16: positive control (DNA extracted from laboratory-bred *Lu. longipalpis* females); 17: positive control (DNA from cultivated *L. (V.) braziliensis*).

>11 Trichophoromyia auraensis

GATGCGTCGAAGTACGAGCAGGCCGACAAGATGCAGCGCGAGCGCGTGGAGGCGAAGAACGGCCTGGAGAACTACGCGTACTCGATGAAGAACACGGTCTCCGACACGAACGTGTCCGGCAAGCTGGAGGA

>3 Pressatia sp

CAACCAGGAGGCGTCGAAGGAAGAGCAGGCCGACAAGATGCAGCGCGAGCGCGTGGAGGCGAAGAACG GCCTGGAGAACTACGCGTACTCGATGAAGAACACGGTCTCCGACACGAACGTGTCCGGCAAGCTGGAC GAGATCGA

>6 Trichophoromyia auraensis

CGTCGAAGGAAGAGCAGGCCGACAAGATGCAGCGCGCGGGGCGTGGAGGACGGCCTGGAGAAC
TACGCGTACTCGATGAAGAACACGGTCTCCGACACGAACGTGTCCGGCAAGCTGGACGAGATCG

>5_Trichophoromyia auraensis

GATGCGTCGAAGTGCGAGCAGGCCGACAAGATGCAGCGCGAGCGCGTGGAGGCGAAGAACGGCCTGGA GAACTACGCGTACTCGATGAAGAACACGGTCTCCGACACGAATGTGTCCGGCAAGCTGGAGGAGAGCG ACAGGTCCGCGCTGAACTCGGCGATCGACACGGCGCTGGAGTGGCTGAAC

>2 Evandromyia saulensis

GATGCGTCGAAGGAAGAGCAGGCCGACAAGGTGCAGCGCGAGCGCGTGGAGGCGAAGAACGGCCTGGA GAACTACGCGTACTCGATGAAGAACACGGTCTCCGACACGAACGTGTCCGGCAAGCTGKACGAGACG

Fig. 2: edited consensus sequences of the 234 bp-hsp70C polymerase chain reaction products.