

<b>Virus Name: Itaituba</b>		<b>Abbreviation: ITAV</b>
Status <b>Possible Arbovirus</b>	Select Agent <b>No</b>	SALS Level <b>3</b>
SALS Basis <b>Insufficient experience with virus; i.e., experience factor from SALS surveys was less than 500 in laboratory facilities with low biocontainment.</b>		
Other Information		
Antigenic Group <b>Phlebotomus Fever</b>		

**SECTION I - Full Virus Name and Prototype Number**

Prototype Strain Number / Designation <b>BeAn 213452</b>	Accession Number	Original Date Submitted <b>9/28/1984</b>
Family <b>Bunyaviridae</b>	Genus <b>Phlebovirus</b>	
Information From <b>F. Pinheiro and A.P.A.T. Rosa</b>	Address <b>Instituto Evandro Chagas, FSESP, Brazilian Ministry of Health, P.O. Box 621, 66.000, Belem, Para, Brazil</b>	
Information Footnote <b>Revised</b>		

**Section II - Original Source**

Isolated By (name) <b>Pinheiro, F.P. et al.</b>	Isolated at Institute <b>Instituto Evandro Chagas, Belem, Para, Brazil</b>	
Host Genus <b>Didelphis marsupialis</b>	Species	Host Age/Stage <b>Young adult</b>
Sex <b>Male</b>		
<u>Isolated From</u> <b>Whole Blood</b>	<u>Isolation Details</u>	
Signs and Symptoms of Illness	Arthropod	
Time Held Alive before Inoculation		
Collection Method <b>UNKNOWN</b>	Collection Date <b>12/7/1971</b>	
Place Collected (Minimum of City, State, Country) <b>Transamazon Highway, Itaituba, Para, Brazil</b>		
Latitude <b>4° 22' S</b>	Longitude <b>55° 50' W</b>	
Macrohabitat <b>Virgin tropical forest</b>	Microhabitat <b>Ground level</b>	Method of Storage until Inoculated <b>-60dC</b>
Footnotes		

**Section III - Method of Isolation**

Inoculation Date  
**1/7/1972**

Animal (Details will be in Section 6)  
**nb mice**

Route Inoculated <b>Intracerebral</b>	Reisolation <b>Not tried</b>
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Other Reasons  
**Virus different from any available in laboratory**

Homologous Antibody Formation by Source Animal  
**Not tested**

Test(s) Used

Footnotes

**Section IV - Virus Properties**

**Physicochemical**

Pieces (number of genome segments)	Infectivity	Sedimentation Coefficients(s) (S)
Percentage wt, of Virion Protein	Lipid	Carbohydrate
Virion Polypeptides: Number	Details	
Non-virion Polypeptides: Number	Details	
Virion Density	Sedimentation Coefficients(s) (S)	
Nucleocapsid Density	Sedimentation Coefficients(s) (S)	

**Stability of Infectivity (effects)**

pH (infective range)

Lipid Solvent (ether - % used to test)	After Treatment Titer	Control Titer
Lipid Solvent (chloroform)	After Treatment Titer	Control Titer
Lipid Solvent (deoxycholate) <b>1:1000</b>	After Treatment Titer <b>&lt;2.5 dex</b>	Control Titer <b>5.2 dex</b>
Other (formalin, radiation)		

**Virion Morphology**

Shape	Dimensions	
Mean nm	Range nm	
Measurement Method	Surface Projections/Envelope	Nucleocapsid Dimensions, Symmetry

**Morphogenesis**

Site of Constituent Formation in Cell

Site of Virion Assembly

Site of Virion Accumulation

Inclusion Bodies

Other

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**Hemagglutination**

Hemagglutination

Antigen Source

Erythrocytes (species used)

**No**

**Mouse and hamster sera tr. by acetone.**

**Goose**

pH Range

pH Optimum

**5.8-7.0**

Temperature Range

Temperature Optimum

**RT and 37dC**

Remarks

Serologic Methods Recommended

**CF, NT**

Footnotes

The virus antigen (mouse brain and liver) did not react in CF testing with the following ascitic fluids: groups A, B, C, Guama, Capim, Bunyamwera; and specific ascitic fluids to Piry, Itaporanga, Jurona, Pacui, and Irituia viruses. Liver antigen gave a reaction of 4/>8 with a grouping ascitic fluid for the Phlebotomus fever group. In a second CF test, this liver antigen was negative with ascitic fluids of viruses of the Phlebotomus fever group: Icoaraci, Bujaru, Anhangá, Sicilian, Chagres, and Urucuri, but reacted with the ascitic fluid for Candiru virus.

Antigens	Ascitic fluids		
	BeAn 213452	Candiru	Control
BeAn 213452	128/1024 <sup>a</sup>	64/256	0
Candiru	128/128	128/256	0
Control	0	0	0

<sup>a</sup> Serum titer/antigen titer; 0 = <4/<4

A cross NT with Candiru gave the following results:

Ascitic fluid	Virus	
	BeAn 213452	Candiru
BeAn 213452	> 5.8 <sup>b</sup>	0.6
Candiru	0.7	3.4
Control	<1.0	0

<sup>b</sup> LNI in dex.

At YARU, it was shown by CF testing that the registered virus is not related to Naples, Sicilian, Ib 15736, Chagres, Punta Toro, and Karimabad viruses. A relationship to Candiru and Nique viruses was demonstrated.

Immune serum to BeAn 213452 (titer of >256) did not react in CF test with the antigens of Anhangá, Bujaru, Chagres, Karimabad, Sicilian, Naples, Arumowot, Icoaraci, Punta Toro, and Gabek Forest. It reacted at 16/8 with Buenaventura and 16/>256 with Nique antigen.

Plaque reduction neutralization tests performed at the National Institute of Health, Hawaii [1] revealed that BeAn 213452 was not neutralized by a 1:10 dilution of the immune sera to the following Phlebotomus fever viruses: Alenquer (BeH 301101), St. Floris, Gordil, Naples, Sicilian, Arumowot, Gabek Forest, Salehabad, Karimabad, Punta Toro, Chagres, Itaporanga, Icoaraci, Candiru, Pacui, Anhangá, Urucuri, Buenaventura, Frijoles, Aguacate, Cacao, Nique, Caimito, Chilibre, Rio Grande, Charleville.

**Section VI - Biologic Characteristics**

Virus Source (all VERTEBRATE isolates)  
**Blood (LV)**

Lab Methods of Virus Recovery (ALL ISOLATIONS)  
**Newborn mice**

Cell system (a)	Virus passage history (b)	Evidence of Infection							Growth Without CPE +/- (g)
		CPE			PLAQUES				
		Day (c)	Extent (d)	Titer TCD50/ml (e)	Day (c)	Size (f)	Titer PFU/ml (e)		
Vero (CL)	SM 7	4	4+	4.5*					

\* Expressed in dex

Vertebrate (species and organ) and arthropod	No. isolations/No. tested	No. with antibody/No. tested Test used	Country and region
Didelphis m. marsupialis (blood)	1		Itaituba, Para, Brazil
Didelphis m. marsupialis		1/38 NT	Itaituba and Altamira, Para, Brazil
Philander o. opossum		0/17 NT	
Marmosa spp.		0/3 NT	
Monodelphis spp.		0/3 NT	
Metachirus nudicaudatus		0/3 NT	
Caluromys spp.		0/3 NT	

No other isolations of the virus were obtained from viscera and/or blood from 921 wild birds, 55 marsupials, 168 rodents, 58 monkeys, 2 edentates, and 2 other animals captured in the Tapacurazinho area in 1971-72. Virus was not isolated from the blood and/or viscera and salivary glands taken from 58 bats collected 25 km. east of Itaituba.

**Section VIII - Susceptibility to Experimental Infection (include viremia)**

Experimental host and age	Passage history and strain	Inoculation Route-Dose	Evidence of infection	AST (days)	Titer log <sub>10</sub> /ml
Mice (nb)	SM 4	ic 0.02	Death	3.5	4.2(brain)  8.8(serum) 8.2(liver)
Mice (nb)		ip 0.02	Death	3.5	
Mice (nb)	SM 5	ic	Death		
Mice (wn)	SM 4	ic 0.03	Antibodies		
Mice (wn)		ip 0.03	Antibodies		
Mice (nb)	SM 3	ic	Viremia, death		
Mice (nb)		ic	Death		
hamsters (23 day)	SM 5	ic	Antibodies		
albino rat		ic	Antibodies		

**Section IX - Experimental Arthropod Infection and Transmission**

Arthropod species & virus source(a)	Method of Infection log <sub>10</sub> /ml (b)		Incubation period (c)		Transmission by bite (d)		Assay of arthropod, log <sub>10</sub> /ml (e)		
	Feeding	Injected	Days	°C	Host	Ratio	Whole	Organ	System

**Section X - Histopathology**

Character of lesions (specify host)	
<u>Inclusion Bodies</u>	<u>Intranuclear</u>
Organs/Tissues Affected	
Category of tropism	

**Section XI - Human Disease**

In Nature	Residual	Death
Subclinical	Overt Disease	
Clinical Manifestations		
Number of Cases	Category (i.e. febrile illness, etc.)	

**Section XII - Geographic Distribution**

Known (Virus detected) <b>Para, Brazil</b>
Suspected (Antibody only detected)

**Section XIII - References**

1. Tesh, R.B. Personal communication.
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**Remarks**

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