

<b>Virus Name: Buenaventura</b>		<b>Abbreviation: BUEV</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>CoAr 3319</b>	Accession Number	Original Date Submitted <b>7/6/1984</b>
Family <b>Bunyaviridae</b>	Genus <b>Phlebovirus</b>	
Information From <b>Robert B. Tesh</b>	Address <b>Pacific Research Section P.O. Box 1680, Honolulu, Hawaii 96806</b>	
Information Footnote <b>Revised</b>		

**Section II - Original Source**

Isolated By (name) <b>Carlos Sanmartin</b>	Isolated at Institute <b>Universidad Del Valle, Cali, Colombia</b>	
Host Genus <b>Lutzomyia (mixed species; pool of 337)</b>	Species	Host Age/Stage <b>Adults</b>
Sex <b>Female</b>		
<u>Isolated From</u>	<u>Isolation Details</u>	
Signs and Symptoms of Illness	Arthropod	
Time Held Alive before Inoculation <b>12-24 hours</b>		
Collection Method <b>Human bait (2)</b>	Collection Date <b>9/1/1964</b>	
Place Collected (Minimum of City, State, Country) <b>Rio Raposo, near Buenaventura, Colombia</b>		
Latitude <b>3° 54' N</b>	Longitude <b>77° 2' W</b>	
Macrohabitat <b>Tropical rain forest covering low hills (&lt;50 meters elevation)</b>	Microhabitat <b>Ground level at early evening (2)</b>	Method of Storage until Inoculated <b>Transported on dry ice; then mechanical freezer at -70dC</b>
Footnotes		

**Section III - Method of Isolation**

Inoculation Date  
**10/15/1964**

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

Route Inoculated <b>Intracerebral</b>	Reisolation <b>Yes</b>
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Other Reasons  
**No other Phlebotomus fever group viruses in laboratory (2)**

Homologous Antibody Formation by Source Animal

Test(s) Used

Footnotes

**Section IV - Virus Properties**

Physicochemical  
**RNA, Single Strand**

Pieces (number of genome segments) <b>3</b>	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)	After Treatment Titer	Control Titer
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)

pH Range

pH Optimum

Temperature Range

Temperature Optimum

Remarks

Serologic Methods Recommended  
CF, NT

Footnotes

**Section V - Antigenic Relationship and Lack of Relationship to Other Viruses**

Buenaventura antigen and hyperimmune serum (homologous titer = 64/512) did not react in cross-CF tests with the following antigens and immune sera: Frijoles, Caimito, Nique, Aguacate, Cacao, Chagres, Candiru, Itaporanga, Urucuri, Pacui, Anhangá, Bujaru, Arumowot, Sicilian, Naples, Gabek Forest, Karimabad, Salehabad, Alenquer, Itaituba, and Charleville [1].

Cross-neutralization tests (plaque method) using Buenaventura virus and hyperimmune serum (homologous titer = 128) were done against each of the following viruses and specific antisera with negative results: Frijoles, Caimito, Nique, Aguacate, Chilibre, Cacao, Chagres, Candiru, Itaporanga, Urucuri, Pacui, Anhangá, Bujaru, Arumowot, Sicilian, Naples, Gabek Forest, Karimabad, Salehabad, Gordil, Saint Floris, Rio Grande, Alenquer, Itaituba and Charleville [1].

Antigenic relationships between Buenaventura, Punta Toro and Icoaraci by both CF and NT were noted as follows [1]:

Antiserum	Antigen/Virus					
	Buenaventura		Punta Toro		Icoaraci	
	CF	NT	CF	NT	CF	NT
Buenaventura	64/512	128	<4/<4	32	<4/<4	<16
Punta Toro	8/32	256	64/256	8192	<4/<4	64
Icoaraci	8/16	32	<4/<4	1024	>2048/256	32000

CF: Antibody titer/antigen titer

NT: Reciprocal of highest serum dilution producing >90% plaque reduction

Buenaventura hyperimmune serum (homologous titer not determined) was tested by HI against 9 other PHL group antigens with the following results: Frijoles (<10), Aguacate (20), Chagres (<10), Icoaraci (80), Itaporanga (10), Anhangá (<10), Arumowot (10), Sicilian (<10) and Karimabad (<10) [1].

**Section VI - Biologic Characteristics**

Virus Source (all VERTEBRATE isolates)

Lab Methods of Virus Recovery (ALL ISOLATIONS)  
Newborn mice and Vero cell cultures

Cell system (a)	Virus passage history (b)	Evidence of Infection						
		CPE			PLAQUES			Growth Without CPE +/- (g)
		Day (c)	Extent (d)	Titer TCD50/ml (e)	Day (c)	Size (f)	Titer PFU/ml (e)	
Vero (CL)	SM 8, Vero 3	5	3-4+		6	pinpoint	5.6*	

\* Expressed in dex

**Section VII - Natural Host Range (Additional text can be added below table)**

Vertebrate (species and organ) and arthropod	No. isolations/No. tested	No. with antibody/No. tested Test used	Country and region
Lutzomyia (mixed species and sexes)	5/about 34,000 *		Raposo River, Buenaventura, Colombia (2)
Lutzomyia (mixed species and sexes)	1/about 150,000 *		Palmares del Pacifico Buenaventura, Colombia (2)

\* See XIV.(1) Remarks

**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)	Prototype; SM 4	ic 0.02	Death (2)	6.6	
Mice (nb)		ip 0.02	None (2)		
Mice (nb)		sc			
Mice (wn)		ic 0.02	None (2)		
Mice (wn)		ip 0.02	None (2)		
Mice (nb)	Prototype; SM 8 Vero 4	ic 0.02	Death (no viremia)	5.8	7.2 (brain)

**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
Aedes albopictus		5.0	10	32			X		No virus present after 10 days as assayed in Vero cells (4).
Culex quinquefasciatus		5.0	10	32			X		Same

**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>Colombia and Panama (2,3)</b>
Suspected (Antibody only detected) <b>1. Tesh, R.B. et al. 1975. Am. J. Trop. Med. Hyg. 24:135-144.</b> <b>2. Sanmartin, C. Personal communication. 1978.</b> <b>3. Tesh, R.B. et al. 1974. Am. J. Trop. Med. Hyg. 23:258-269.</b> <b>4. Tesh, R.B. 1975. J. Med. Ent. 12:1-4.</b>

**Section XIII - References**

<b>1. Tesh, R.B. et al. 1975. Am. J. Trop. Med. Hyg. 24:135-144.</b> <b>2. Sanmartin, C. Personal communication. 1978.</b> <b>3. Tesh, R.B. et al. 1974. Am. J. Trop. Med. Hyg. 23:258-269.</b> <b>4. Tesh, R.B. 1975. J. Med. Ent. 12:1-4.</b>
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**Remarks**

<b>Several closely related or identical viruses isolated from Lutzomyia (mixed species and sexes) in Panama (3).</b>
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