

Virus Name: Arbia		Abbreviation: ARBV
Status <b>Probable 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>ISS.Phl.18</b>	Accession Number	Original Date Submitted <b>5/1/1985</b>
Family <b>Bunyaviridae</b>	Genus <b>Phlebovirus</b>	
Information From <b>P. Verani, et. al.</b>	Address <b>Istituto Superiore di Sanita, Viale Regina Elena 299, I-00161 Rome, Italy</b>	
Information Footnote <b>Reviewed by editor</b>		

#### Section II - Original Source

Section 11 - Original Source		
Isolated By (name) <b>P. Verani, et. al. (1,6)</b>	Isolated at Institute <b>Istituto Superiore di Sanita</b>	
Host Genus <b>Phlebotomus perniciosus</b>	Species	Host Age/Stage
Sex <b>Female</b>		
<u>Isolated From</u>	<u>Isolation Details</u>	
Signs and Symptoms of Illness	Arthropod <b>Engorged</b>	
Time Held Alive before Inoculation <b>A few hours</b>		
Collection Method <b>Caught by hand</b>	Collection Date <b>7/25/1980</b>	
Place Collected (Minimum of City, State, Country) <b>Sesto Fiorentino, Toscana, Italy</b>		
Latitude <b>43° 50' N</b>	Longitude <b>11° 10' E</b>	
Macrohabitat <b>Hills covered by deciduous woods (250 meters alt.)</b>	Microhabitat <b>Poultry yard</b>	Method of Storage until Inoculated <b>-70dC</b>
Footnotes		

### Section III - Method of Isolation

Inoculation Date  
**10/3/1980**

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

Route Inoculated  
**Intracerebral**

Reisolation

Other Reasons

Homologous Antibody Formation by Source Animal

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)	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)

**Yes**

**SMB ext. by sucrose-acetone**

**Goose**

pH Range

pH Optimum

**5.75-6.2**

**6.0**

Temperature Range

Temperature Optimum

**22dC-37dC**

**37dC**

Remarks

**HA production is difficult, usually of poor titre**

Serologic Methods Recommended

**CF, NT, IFA**

Footnotes

**HA production is difficult, usually of poor titre**

ISS.Phl.18 antigen and MIAF did not react in CF test with antigens and MIAF prepared against Toscana , SFN, and SFS viruses.

At YARU [2] ; ISS.Phl.18 was screened by indirect fluorescent antibody test against the following MIAF: Phlebotomus fever group, Toscana , SFN, Arumowot, Gordil, Saint Floris, SFS, Rift Valley Fever , Karimabad, Salehabad, Gabek Forest, VSV group, Isfahan , Chandipura , and Jug Bogdanovac. Positive reactions were obtained with Toscana , SFN, Gordil, Saint Floris, Salehabad, Gabek Forest, and the Phlebotomus fever group MIAF. ISS.Phl.18 antigen and MIAF were then tested by CF against Salehabad, Gabek Forest, Gordil, Toscana , SFN, Saint Floris antigens, and MIAF. Positive reactions were obtained between ISS.Phl.18, Salehabad, and Gabek Forest.

ANTIGEN	MIAF		
	ISS.Phl.18	Salehabad	Gabek Forest
ISS.Phl.18	256/> 256 <sup>a</sup>	128/>64	<4/<4
Salehabad	64/> 128	256/> 128	<4/<4
Gabek Forest	16/4	<4/<4	>256/> 128
<sup>a</sup> Serum titer/antigen titer.			

Cross NT tests (plaque reduction method) between ISS.Phl.18 and Salehabad viruses and MIAF gave the following results:

VIRUS	MIAF	
	ISS.Phl.18	Salehabad
ISS.Phl.18	160 <sup>b</sup>	<10
Salehabad	<10	320
<sup>b</sup> Dilution producing >90% plaque reduction.		

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)	SMB 4	3	4+	6.4 °	5	pinpoint	7.6°	
Vero (CL)	Vero 3	3	4+		5	pinpoint	5.9	
° Expressed in dex								

Vertebrate (species and organ) and arthropod	No. isolations/No. tested	No. with antibody/No. tsted Test used	Country and region
Man		0/105 HI	Toscana, Italy
Chicken		0/65 HI	
Rabbit		0/2 HI	
Phlebotomus perniciosus	3/609		Florence province, Toscana,Italy(1,6)
Phlebotomus perniciosus(males)	1		
Phlebotomus perfiliewi	23/9,351		Siena province,Toscana, Italy (3)
Phlebotomus perfiliewi (males)	3		
Phlebotomus perfiliewi	2/882		Teramo province, Abruzzo, Italy

Experimental host and age	Passage history and strain	Inoculation Route-Dose	Evidence of infection	AST (days)	Titer log10/ml	
Mice (nb)	SMB5	ic 0.01	Death	4	8.24	
Mice (nb)		ip				
Mice (nb)		sc				
Mice (wn)	SMB 6	ic 0.03	Death	6		
Mice (wn)	SMB 4	ip 0.05	Antibodies			
rabbit		ic 0.20	Antibodies			
rabbit		ip 1.00	Antibodies			

# Section IX - Experimental Arthropod Infection and Transmission

Arthropod species & virus source(a)	Method of Infection log10/ml (b)		Incubation period (c)		Transmission by bite (d)		Assay of arthropod, log10/ml (e)		
	Feeding	Injected	Days	°C	Host	Ratio	Whole	Organ	System
Phlebotomus perniciosus , artificial "blood" meal	8.8		8-10	26 °			3.5 *		Vero Cells (TCID50)
Arbia virus replicated in intrathoracically inoculated female Phlebotomus perniciosus . Sandflies were refractory or resistant to oral infection (4,5). Experimental transovarialtransmission of Arbia virus demonstrated (5).									

\* Mean virus titer per insect for 10 positive female insects.

## Section X - Histopathology

Character of lesions (specify host)

Inclusion Bodies

Intranuclear

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)

**Central Italy**

Suspected (Antibody only detected)

**Section XIII - References**

1. Verani, P. et al. 1982. Unpublished data.
2. Tesh, R.B. Personal communication. 1983.
3. Verani, P. et al. 1983. Unpublished data.
4. Ciufolini, M.G., et al. 1985. Am. J. Trop. Med. Hyg. 34:174-179.
5. Tesh, R.B. and Modi, G.B. 1984. Am. J. Trop. Med. Hyg. 33:1007-1016.
6. Verani, P., et al. 1988. Am. J. Trop. Med. Hyg. 38:433-439.

**Remarks**