Virus Name: Rift Valley fever Abbreviation: RVFV

Status Select SALS Level

Arbovirus Agent 3

Yes

SALS Basis

Results of SALS surveys and information from the Catalogue.

Other Information

USDA Permit Required, DOC Permit Required, Hepa Filtration, USDA Restricted, USDA High

Consequence Agent, Vaccination Recommended

Antigenic Group Phlebotomus Fever

SECTION I - Full Virus Name and Prototype Number

Prototype Strain Number / Designation Accession Number Original Date Submitted

2/3/1985

Family Genus
Bunyaviridae Phlebovirus

Information From Address

M.P. Weinbren Puerto Rico Nuclear Center, Caparra Heights Station, San Juan, P.R.

Information Footnote Reviewed by editor

Section II - Original Source

Isolated By (name) Isolated at Institute

Daubney, et al. (1) Vet. Res. Lab., Kabete, Kenya

Host Genus Species Host Age/Stage
Sheep Lamb (newborn)

Sex

Not Answered

<u>Isolated From</u> <u>Isolation Details</u>

Whole Blood

Signs and Symptoms of Illness Arthropod

Abortion and death in sheep, especially high mortality

in young lambs.*

Time Held Alive before Inoculation

Collection Method Collection Date Not stated 7/27/1930

Place Collected (Minimum of City, State, Country)

North of Lake Naivasha, Kenya

Latitude Longitude 0° 44' S 36° 26' E

Macrohabitat Microhabitat Method of Storage until

Upland tropical savannah Daylight, ground level, outdoors Inoculated

Not stated

Footnotes

Section III - Method of Isolation

Inoculation Date 7/28/1930

Animal (Details will be in Section 6)

Sheep

Route Inoculated Reisolation Intravenous Yes

Other Reasons

Other isolations of identical virus in same outbreak

Homologous Antibody Formation by Source Animal

Test(s) Used

Footnotes

Section IV - Virus Properties

Physicochemical

RNA

Pieces (number of genome

Infectivity

Sedimentation Coefficients(s)

(S)

segments) 3 (22)

Percentage wt, of Virion Protein

Lipid

Carbohydrate

Virion Polypeptides: Number

Details

3

Non-virion Polypeptides: Number Details

Virion Density

Sedimentation Coefficients(s)

Nucleocapsid Density

Sedimentation Coefficients(s)

Stability of Infectivity (effects)

pH (infective range)

Lipid Solvent (ether - % used to

After Treatment Titer

test)

<2.0 dex

Control Titer

4.0 dex (9)

Lipid Solvent (chloroform)

After Treatment Titer

Control Titer

Lipid Solvent (deoxycholate)

After Treatment Titer 2.0 dex less (6)

Control Titer

Other (formalin, radiation)

Inact. by 0.25% formalin, 4C/3 days; by methylene blue+light (15).

Virion Morphology

Shape Dimensions

94-100 nm (20-22)

Mean

Range

Measurement Method Surface Projections/Envelope Nucleocapsid Dimensions,

Electron microscopy (22) Envelope observed (22) Symmetri

Morphogenesis

Site of Constituent Formation in

Site of Virion Assembly

Site of Virion Accumulation

Cell

Inclusion Bodies

Other

Hemagglutination

Hemaggiutination Antigen Source Erythrocytes (species used)

Yes Mouse serum, acetone-ether; freezing 72 hours (7); Go

Goose

sucrose-acetone (8)

pH Range Varies pH Optimum

Temperature Range

Temperature Optimum

4-37dC 25dC

Remarks

Frozen serum of naturally infected animal has been successfully used as antigen in HI test for diagnosis (11).* Short illness, loss of appetite, listlessness, asthenia.

Serologic Methods Recommended

HI, CF, NT

Footnotes

Frozen serum of naturally infected animal has been successfully used as antigen in HI test for diagnosis (11).* Short illness, loss of appetite, listlessness, asthenia.

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

RVF virus antigenically related to viruses of the PHL serogroup by HI and plaque-reduction neutralization tests [23].

Zinga virus has been shown to be identical to RVF virus by IFA and NT [30], [31].

Section VI - Biologic Characteristics

Virus Source (all VERTEBRATE isolates)
Blood (M) (LV), liver (LV), spleen (LV), milk (LV), urine (LV),
feces (LV)

Lab Methods of Virus Recovery (ALL ISOLATIONS)

Newborn and weanling mice, baby chick, chick embryo, hamster, primates

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

Rift Valley fever virus grows readily in virtually all types of cell culture.

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
Man (many isolations from veterinarians and lab workers infected while handling animals, tissues or virus).	1	(Naturally infected case)	South Africa (10)
Man		146/2,223 CF	Nigeria (26)
Man	7		Uganda (19)
Lambs, sheep, cattle	Many		South Africa (1, 3, 4, 10, 11)
Man (serum)	10		Egypt (24)
Man (serum)	53		Egypt (25)
Wild rodents		0/106 NT	Uganda (19)
Mosquitoes of various species-numerous isolations, likely vectors:			
Eretmapodites chrysogaster group(2) Aedes (Ochlerotatus) caballus(4) Aedes (Neomelaniconion) circumluteolus(5) Culex (Culex) theileri(11)			
Culicoides spp.	1		Nigeria (32)
Micropteropus pusillus (bat)	1		Guinea (34)
Hipposideros abae (bat)	1		Guinea (34)

Primary hosts probably are sheep, cattle, buffalo, certain antelopes and rodents (1,3,4,10,11) all infected by mosquitoes; and man - most commonly infected while handlinng sick or dead animals, but sometimes by mosquitoes. Epizootics have caused enormous economic losses. Culex (Culex) theileri probably main epizootic vector (33).

Section VIII - Susceptibility to Experimental Infection (include viremia)

Experimental host and age	Passage history and strain	Inoculation Route-Dose	Evidence of infection	(days)	Titer log10/ml
Mice (nb)	Uganda	ic 0.03	Death	1-4	9.0
Mice (nb)		ip 0.03	Death	2-6	8.0
Mice (nb)		sc			
Mice (wn)		ic 0.03	Death	2-5	8.0
Mice (wn)		ip 0.06	Death	1-7	6.6
rhesus monkey (ad)		sc	Mild febrile or asymptomatic infection with resulting antibody response		
Cercopithecus aethiops (ad)		sc	Same as rhesus		
newborn merino lambs		sc	Death	1-2	
Pregnant ewes		sc	Abortion, fever, sometimes death		
Field voles, dormice, wood mice			Susceptible (12)		
kittens, puppies		sc	Death (18)		
5 African buffalo (Syncerus caffer)			1/2 pregnant aborted; 4/5 48 hr viremic (29)		

Section IX - Experimental Arthropod Infection and Transmission Arthropod species Method of Infection Incubation Transmision by Assay of arthropod, log10/ml & virus source(a) log10/ml (b) period (c) bite (d) (e) °C Ratio Feeding Injected Host Whole Organ System Days Virus has been transmitted experimentally by the bite of Erchrysogaster(13), Ae caballus (4), Ae aegypti (8). The first two of these and other mosquitoes are probably natural vectors. Culex pipiens: Laboratory reared mosquitoes from Nile Delta epizootic area transmitted RVF virus following feeding on viremic hamsters. Infection rata = 87%; trans. rate = 40% (28).

Section X - Histopathology

Character of lesions (specify host)

Lambs, mice: generalized hepatic necrosis with intranuclear acidophilic histpath inclusions. Nuclear chromatin retains basophilia - distinguishing them from yellow fever. Hyaline acidophilic bodies in cytoplams in RVF - while in YF the whole cell is acidophilic and there are no cytoplasmic bodies.

Inclusion Bodies

<u>Intranuclear</u>

Lower Vertabrates

Organs/Tissues Affected Brain (LV)*, liver (LV).

Category of tropism

* Sheep, goats, hepatotropic. Mice convert to neurotropic on serial ic inoculation

Section XI - Human Disease

In Nature Residual Death Significant Significant Reported

Subclinical Overt Disease Significant Significant

Clinical Manifestations

Fever, headache, prostration, conjunctival inflammation, stiff nect, myalgia, arthralgia, CNS signs (including encephalitis, hemorrhagic signs, lymphadenopathy, vomiting, central scotoma- detached retina (1,16,17).

Number of Cases

Category (i.e. febrile illness, etc.)

300 cases in detail; numerous others reported (8). Febrile illness

Section XII - Geographic Distribution

Known (Virus detected)

Kenya, Uganda, South Africa (1,2,3,4,5,11), Egypt (24,25)

Suspected (Antibody only detected)

Sudan, Nigeria (26)

Section XIII - References

- DAUBNEY, R., et al. 1931. Jour. Path. and Bact. 34:545-579.
- SMITHBURN, K.C., et al. 1948. Brit. J. Exp. Path. 29:107-121.
- ALEXANDER, R.A. 1951. J. S. Afr. Vet. Med. Assn. 22:105.
- GEAR, J., et al. 1955. S. Afr. Med. J. 29:514.
- KOKERNOT, R.H., et al. 1957. S. Afr. J. Med. Sci. 22:71-80.
- THEILER, M. 1957. Proc. Soc. Exp. Biol. and Med. 96:380-382.
- MIMS, C.A.C. and MASON, P.J. 1956. Brit. J. Exp. Path. 37:423-433.
- 8. WEINBREN, M.P. Doctoral Thesis, Witwatersrand Univ. Medical School.
- 9. ANDREWES, C.H. and HORSTMAN, D. 1949. J. Gen. Microbiol. 3:290-297.
- 10. GEAR, J., et al. 1951. S. Afr. Med. J. 25:908-912.
- 11. SMITHBURN, K.C.Unpublished results.
- 12. FINDLAY, G.M. 1931. Trans. Roy. Soc. Trop. Med. and Hyg. 25:229.
- 13. SMITHBURN, K.C., et al. 1949. Brit. J. Exp. Path. 30:35-47.
- 14. KOKERNOT, R.H., et al. 1956. J. Immunol. 77:313-323.
- 15. MACKENZIE, R.D. 1935. J. Path. and Bact. 40:65.
- 16. SCHRIRE, L. 1951. S.Afr. Med. J. 25:926.
- 17. FREED, I. 1951. S. Afr. Med. J. 25:930.
- 18. MITTEN, J.G., et al. 1970. J. Infect. Dis. 121:25-31.
- 19. HENDERSON, B.E., et al. 1972. Ann. Trop. Med. Parasit. 66:343-355.
- Lecastas, G. and WEISS, K.W. 1968. Arch. ges. Virusforsch. 25:58-64.
- 21. POLSON, A. and STANDARD L. 1970. Virology 40:781-791.
- MURPHY, F.A. 1973. Intervirology 1:297-316.
- 23. SHOPE, R.E., et al. 1980. Lancet. Vol. 1 886-888.
- ABDEL-WAHAB, K.S.E.D., et al. 1978. Trans. Roy. Soc. Trop. Med. Hyg. 72:392-396.
- 25. IMAM, I.Z.E. 1979. Bull. Wld. Hlth. Org. 57:100%-439.
- 26. TOMORI, O. 1980. J. Med. Virol. 5:343-350.
- 27. RICE, R.M., et al. 1980. Virology 105:256-260.
- 28. MEEGAN, J.M., et al. 1980. Am. J. Trop. Med. Hyg. 29:1405-1410.
- DAVIES, P.G. and KARSTAD, L. 1981. Trop. Anim. Hlth Prod. 13:185-188.
- 30. SHOPE, R.E. Personal communication, 1982.
- 31. MEEGAN, J.M. 1983. Lancet i: 641.
- 32. FABIYI, A. 1980. Zbl. Bakt. Suppl 9:215-218.
- 33. MCINTOSH, B.M. 1975. Unpublished.
- 34. BOIRO, I., et al. 1987. Bull. Soc. Path. Ex. 80:62-67.

Remarks

Many infections in veterinary field officers and laboratory workers - often with prolonged convalescence; eye complications common in man; herdsmen often become infected. Neuro-adapted virus used with success in South African Veterinary Service as a living virus vaccine. Dead virus vaccine recently prepared in U.S. evokes antibody response.