

<b>Virus Name: Kunjin</b>		<b>Abbreviation: KUNV</b>
Status <b>Arbovirus</b>	Select Agent <b>No</b>	SALS Level <b>2</b>
SALS Basis <b>Results of SALS surveys and information from the Catalogue.</b>		
Other Information		
Antigenic Group <b>B</b>		

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

Prototype Strain Number / Designation <b>MRM16</b>	Accession Number	Original Date Submitted <b>1/5/1985</b>
Family <b>Flaviviridae</b>	Genus <b>Flavivirus</b>	
Information From <b>R.L. Doherty</b>	Address <b>Queensland Institute of Medical Research, Herston Rd., Herston, Brisbane</b>	
Information Footnote <b>Reviewed by editor</b>		

**Section II - Original Source**

Isolated By (name) <b>Doherty, et al. (1)</b>	Isolated at Institute <b>Brisbane</b>	
Host Genus <b>Culex annulirostris Skuse</b>	Species	Host Age/Stage <b>Adult</b>
Sex <b>Female</b>		
<u>Isolated From</u>	<u>Isolation Details</u>	
Signs and Symptoms of Illness	Arthropod	
Time Held Alive before Inoculation		
Collection Method <b>Aspirated from horses or man</b>	Collection Date <b>4/2/1960</b>	
Place Collected (Minimum of City, State, Country) <b>Mitchell River Mission, Queensland, AS</b>		
Latitude <b>15° 30' S</b>	Longitude <b>141° 40' E</b>	
Macrohabitat <b>Low-lying plain bordering Gulf of Carpentaria; open forest-grassland</b>	Microhabitat <b>On bank of creek at edge of aboriginal mission</b>	Method of Storage until Inoculated <b>Dry ice (-70dC) and Revco at -60dC</b>
Footnotes		

**Section III - Method of Isolation**

Inoculation Date  
**4/26/1960**

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

Route Inoculated <b>Intracerebral</b>	Reisolation <b>Yes</b>
--	---------------------------

Other Reasons  
**Subsequent isolations and demonstration of antibody in the same area.**

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 <b>Electrophoresis on polyacrylamide gel showed four proteins in virion and at least three other virus-coded proteins in infected cells (15).</b>	
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	

**Hemagglutination**

Hemagglutination <b>Yes</b>	Antigen Source <b>SMB ext. by sucrose-acetone or acetone-ether</b>	Erythrocytes (species used) <b>Goose</b>
--------------------------------	---	---

pH Range <b>6.4-7.3</b>	pH Optimum <b>6.6</b>
----------------------------	--------------------------

Temperature Range	Temperature Optimum <b>37dC used routinely</b>
-------------------	---

Remarks

Serologic Methods Recommended  
**CF, NT**

Footnotes

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

Immune Sera (mouse)	MRM16 Antigen			Antigens	MRM16 Mouse Antiserum		
	HI Ht/Ho	CF Ht/Ho	NT Ht/Ho		HIT Ht/Ho	CF Ht/Ho	NT Ht/Ho
Studies in Brisbane [1]							
MVE	320/640	32/128	<2.0/3.0	MVE	640/320	64/64	2.8/>6.0
Kokobera (MRM32)	40/640	<8/32	<1.5/>5.0	Kokobera	320/320	<8/64	2.2/>6.0
Edge Hill (C281)	40/320	<8/128	<1.5/>3.2	Edge Hill	640/320	128/64	1.7/>6.0
Stratford (C338)	80/320	8/64	3.0/3.0	Stratford	640/320	64/64	2.0/>6.0
Studies at RFVL in New York [2]							
Dengue 4		16/>128		RSSE		0/256	
RSSE	20/320	<4/32		Tembusu		0/128	
Tembusu		4/32		SLE		32/128	
Rio Bravo	320/5120			West Nile		16/128	
Russuquara	20/160			IBF		8/128	

Bussuquara	20/160		SLE	0/120
Modoc	640/5120		Rio Bravo	0/320
Usutu	320/1280		Bussuquara	10/320
Israel Turkey Men.	0/640		Modoc	0/320
Wesselsbron	10/40		Usutu	0/320
Spondweni	20/320		IT	80/320
YF	320/5120		Wesselsbron	0/320
Banzi	160/640		Spondweni	0/320
Zika	160/2560		YF	0/320
Ilheus	1280/5120		Banzi	0/320
JBE	80/2560		Zika	20/320
SLE	320/2560		Ilheus	0/320

NT: LNI given in dex

Studies by plaque reduction (PR) and plaque inhibition (PI) which suggest that Kunjin and West Nile are closely related [3].

Sera (rabbit antisera)	MRM61 Virus		Viruses	MRM61 Rabbit Antiserum	
	PR	PI		PR	PI
MVE	0.73/1.55	8-10/18-19	MVE	0.83/2.76	+/25-26
Edge Hill (C281)	0.32/1.91	0/18-20	Edge Hill	0.47/2.76	0/25-26
West Nile	1.24/3.31	10-13-20/22	West Nile	2.70/2.76	14-16/25-26

PR: LNI given in dex.

PI: Given as zone(s) of inhibition

## 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						
		CPE			PLAQUES			Growth Without CPE +/- (g)
		Day (c)	Extent (d)	Titer TCD50/ml (e)	Day (c)	Size (f)	Titer PFU/ml (e)	
Chick embryo (PC)	MRM 61 SMB 4	3	CPE		3-4	Plaques	8.1*(3)	
PS (CL)	MRM 61 SMB 5	3	4+	8.0*	4	3-4 mm	8.7 (11)	
BHK-21 (CL)	MRM 16 SMB 2	3,4	4+	8.3 (12)				
Vero (CL)	SMB 7				2	12 mm	8.1 (13)	
LLC-MK2 (CL)					2	2 mm	8.3 (13)	
Aedes albopictus(CL)								+ (17)
Ae aegypti(CL)								+ (14)

\* Expressed in dex

Vertebrate (species and organ) and arthropod	No. isolations/No. tested	No. with antibody/No. tested Test used	Country and region
Man	One isolation from laboratory infection		Brisbane, Australia (4)
Domestic fowl		14/73	NW Queensland, AS; 1959 survey (5)
Cattle		6/419 HI	Queensland, AS; 1965 (10)
Cx annulirostris	9/6703		Mitchell River, Mission, AS; 1960
Cx annulirostris	0/2260		Mitchell River, Mission, AS; 1961
Cx annulirostris	5/914		Mitchell River, Mission, AS; 1963
Cx squamosus	1/48		
Cx pseudovishnui group	3/73 pools		Sarawak; 1966 (9) Mitchell River, AS; 1966 (18)
Oriolus flavocinctus	1		
Horses	1/2 *		Australia (19)

\* Virus isolated from cervical spinal cord of 1 of 2 horses both ill with encephalomyelitis; both were euthanized (19).

Cross-reaction between group B viruses makes it difficult to interpret serum surveys in North Queensland. However six human sera reacting to Kunjin only have been found in Cape York Peninsula (5).

Experimental host and age	Passage history and strain	Inoculation Route-Dose	Evidence of infection	AST (days)	Titer log <sub>10</sub> /ml	
Mice (nb)	MRM16, SMB 3	ic 0.01	Death	4	10.0	
Mice (nb)		ip 0.03	Death	5	9.5	
Mice (nb)		sc				
Mice (wn)		ic 0.03	Death	5	8.5	
Mice (wn)		ip 0.03	Few deaths only	10-14		
mice (nb)	MRM61, SMB 4	ic 0.01	Death	4	8.4	
embryonated egg (7 day)	MRM16, SMB 3	ys 0.1	Death	<3	9.3(lesions)	
™ (11 day)		CAM 0.05	Death and pocks		3.5(death)	
rabbit (ad)		ip 0.2	Antibody formation			
guinea pig (ad)		ip 0.2	Antibody formation			
calf (2 mo)	SMB 7, 8	various	Clinical, pathol. and antibody (8)			

**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
Culex quinquefasciatus	Adult females intrathoracically inoculated and membrane fed; virus multiplication demonstrated by titration in mice; 6.8/mosquito at 8 days; 5.5/mosquito at 25 days. Aedes aegypti infected by feeding transmitted virus by bite at 15-25 days (20). Exp. transovarial transmission demonstrated with Aedes albopictus (21).								

**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  
**Reported**

Overt Disease  
**Reported**

Clinical Manifestations  
**Fever (R), rash (R), lymphadenopathy (R)**

Number of Cases  
**1 (22)**

Category (i.e. febrile illness, etc.)  
**Encephalitis (22)**

**Section XII - Geographic Distribution**

Known (Virus detected)

**Australia; Borneo, Indonesia; Sarawak, Malaysia.**

Suspected (Antibody only detected)

**Northern West Australia (7); Sepik District, New Guinea (16)**

### Section XIII - References

1. Doherty, R.L., et al. 1963. *Aust. J. Exp. Biol. Med. Sci.* 41:17-40.
2. Theiler, M. Personal communication. 1961.
3. Westaway, E.G. 1965. *Virology* 26:528-537.
4. Allan, B.C., et al. 1966. *Med. J. Aust.* 2:844-847.
5. Doherty, R.L., et al. 1964. *Aust. J. Exp. Biol. Med. Sci.* 42:149-164.
6. Standfast, H.A. and Carley, J.G. Personal communication. 1963.
7. Stanley, N.F. and Choo, S.B. 1964. *Bull. World Health Organ.* 30:221-226.
8. Spradbrow, P.B. and Clark, L. 1966. *Aust. Vet. J.* 42:65-69.
9. Bowen, E.T.W., et al. 1970. *Ann. Trop. Med. Parasitol.* 64:263-268.
10. Sanderson, C.J. 1969. *Am. J. Trop. Med. Hyg.* 18:433-439.
11. Westaway, E.G. 1966. *Am. J. Epidemiology* 84:439-456.
12. Karabatsos, N. and Buckley, S.M. 1967. *Am. J. Trop. Med. Hyg.* 16:99-105.
13. Stim, T.B. 1969. *J. Gen. Virol.* 5:329-338.
14. Rehacek, J. 1968. *Acta Virol.* 12:241-246.
15. Westaway, E.C. and Reedman, B.M. 1969. *J. Virol.* 4:688-693.
16. Marshall, I. Personal communication. 1971.
17. Singh, K.R.P. 1972. *Adv. Virus Res.* 17:187-206.
18. Whitehead, R.H., et al. 1968. *Trans. R. Soc. Trop. Med. Hyg.* 62:439-445.
19. Badman, R.T., et al. 1984. *Commun. Diseases Intelligence.* No. 84/17. pp. 5-6.
20. Kay, B.H., et al. 1974. Unpublished data.
21. Tesh, R.B. 1980. *Am. J. Trop. Med. Hyg.* 29:657-666.
22. Muller, D., et al. 1986. *Med. J. Australia.* 144:41-42.

### Remarks