

<b>Virus Name: Marburg</b>		<b>Abbreviation: MBGV</b>
Status <b>Probably not Arbovirus</b>	Select Agent <b>Yes</b>	SALS Level <b>4</b>
SALS Basis <b>Results of SALS surveys and information from the Catalogue.</b>		
Other Information <b>DOC Permit Required</b>		
Antigenic Group <b>Marburg</b>		

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

Prototype Strain Number / Designation <b>M.1/2 (1, 3)</b>	Accession Number	Original Date Submitted <b>8/16/1984</b>
Family <b>Not listed</b>	Genus <b>Not listed</b>	
Information From <b>R. Siegert and D.I.H. Simpson</b>	Address <b>Hygiene-Institut de Universitat, 355 Marburg/L, Pilgrimstein 2, and Queen's Univ. of Belfast, N. Ireland</b>	
Information Footnote <b>Reviewed by editor</b>		

**Section II - Original Source**

Isolated By (name) <b>R. Siegert</b>	Isolated at Institute <b>Hygiene-Institut, Marburg</b>	
Host Genus <b>Man</b>	Species	Host Age/Stage <b>40 years</b>
Sex <b>Male</b>		
<u>Isolated From</u> <b>Whole Blood</b>	<u>Isolation Details</u>	
Signs and Symptoms of Illness <b>Several, see XI (3) and XI (4)</b>	Arthropod	
Time Held Alive before Inoculation		
Collection Method <b>Venepuncture</b>	Collection Date <b>8/22/1967</b>	
Place Collected (Minimum of City, State, Country) <b>Marburg/L, Med. Klinik d. Universitat</b>		
Latitude <b>50° 50' N</b>	Longitude <b>8° 45' E</b>	
Macrohabitat	Microhabitat	Method of Storage until Inoculated <b>No storage</b>
Footnotes		

**Section III - Method of Isolation**

Inoculation Date  
**8/22/1967**

Animal (Details will be in Section 6)  
**guinea pig**

Route Inoculated <b>Intraperitoneal</b>	Reisolation <b>Yes</b>
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Other Reasons  
**Electron microscopy(1,7); repeated isolations in other laboratories (11,21)**

Homologous Antibody Formation by Source Animal  
**Yes**

Test(s) Used  
**CF, NT, IF (1,4,5,6)**

Footnotes

**Section IV - Virus Properties**

Physicochemical  
**RNA**

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) <b>1:1</b>	After Treatment Titer <b>0 dex</b>	Control Titer <b>6.0 dex</b>
Lipid Solvent (chloroform) <b>Also used (4)</b>	After Treatment Titer	Control Titer
Lipid Solvent (deoxycholate) <b>1:1000</b>	After Treatment Titer <b>0 dex</b>	Control Titer <b>6.0 dex</b>
Other (formalin, radiation)		

**Virion Morphology**

Shape <b>Rhabdovirus-like morphology</b>	Dimensions <b>70 x 665 nm</b>	
Mean nm	Range nm	
Measurement Method <b>Electron microscopy (7)</b>	Surface Projections/Envelope <b>Envelope observed</b>	Nucleocapsid Dimensions, Symmetry

### Morphogenesis

Site of Constituent Formation in Cell	Site of Virion Assembly	Site of Virion Accumulation
Inclusion Bodies	Other	

### Hemagglutination

Hemagglutination <b>No</b>	Antigen Source <b>Cell culture fluid; infected guinea pig liver, spleen.</b>	Erythrocytes (species used) <b>Several**</b>
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pH Range	pH Optimum
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Temperature Range	Temperature Optimum
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#### Remarks

**CF antigen prepared from "chronically" infected Vero carrier cultures (9, 10) (All organ extracts were nonspecific.) \*\*  
Chicken, guinea pig, goose, mouse, man**

#### Serologic Methods Recommended

**NT, CF**

#### Footnotes

**CF antigen prepared from "chronically" infected Vero carrier cultures (9, 10) (All organ extracts were nonspecific.) \*\*  
Chicken, guinea pig, goose, mouse, man**

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

Marburg virus is an antigenically distinct virus, unrelated to any of almost 200 viruses with which it has been compared [10].

Human convalescent and guinea pig immune sera showed negative results when tested by HI or CF against a large number of arbovirus and non-arbovirus antigens, including rhabdoviruses, Tacaribe group agents, and viruses causing hemorrhagic disease manifestations in man, monkeys or deer [10].

Marburg virus was shown to be distantly related, by IFA, to the subsequently isolated Ebola virus. See Ebola virus registration card.

**Section VI - Biologic Characteristics**

Virus Source (all VERTEBRATE isolates)  
**Blood (M)(LV), CNS (LV), liver (M), spleen (M),  
nasopharyngeal (M), salivary gland (M) (from saliva swab)**

Lab Methods of Virus Recovery (ALL ISOLATIONS)  
**Newborn mice and suckling hamsters; WI-26 human diploid  
embryo fibroblast**

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)		

Marburg virus infects and replicates in a variety of mammalian cell cultures, including primary and continuous cell cultures, primary and continuous primate cell cultures, primary and continuous guinea pig cell cultures and BHK-21 cell cultures. In most instances, the development of CPE was variable or absent. One primary and one continuous primate cell culture, a continuous human cell culture and BHK-21 exhibited distinct CPE after infection with Marburg virus. Variable CPE was produced in Vero cell cultures, and CPE endpoints generally were 10-100 times lower than infective virus titrations. See Reference 14.

Vertebrate (species and organ) and arthropod	No. isolations/No. tested	No. with antibody/No. tested Test used	Country and region
Man	31/many		Marburg; Frankfurt, W. Germany
Man	3		Southern Africa (23, 24)
Man	4		Belgrade, Yugoslavia, West Germany
Cercopithecus aethiops *			No specific CF antibodies found in monkeys (9). Positive CF tests have been reported with sera from primate species as possible evidence of natural infection of primates in Uganda (22), but specificity of CF antibodies employing infected guinea pig crude liver antigens still questionable.
Man		22/22 CF	Marburg (6)
Man		1% / >400 IFA	Liberia (26)
Vervet monkeys		2 IFA	Kenya (25)
Baboons		3 IFA	

\* No isolations from naturally infected animals.

Experimental host and age	Passage history and strain	Inoculation Route-Dose	Evidence of infection	AST (days)	Titer log <sub>10</sub> /ml
Mice (nb)		ic	Inapparent infection (15)		
Mice (nb)		ip	Inapparent infection (15)		
Mice (nb)		sc			
Mice (wn)		ic	Inapparent infection (15)		
Mice (wn)		ip	Inapparent infection (15)		
guinea pigs (ad)	Original human	ic 0.1	Febrile illness only (1,4,8)		
guinea pigs		ip 0.1	Febrile illness only (1,4,8)		
guinea pigs		sc 0.1	Febrile illness only (1,4,8)		
guinea pigs (ad)	GP 1	ic 0.1	Sickness and death	13-15	
guinea pigs		ip 0.1	Sickness and death		
guinea pigs		sc 0.1	Sickness and death		
monkeys (ad)	Original human	sc	Sickness and death (11,12)	6-10	
(rhesus, vervet squirrel)	+GP passage	ip	Sickness and death (11,12)	6-10	
hamster (nb)	GP 9	ic 0.02	Death (16)	8-15	
hamster (nb)	hamster 9	ip 0.02	Death (16)		

**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

Marburg virus found to replicate in *Aedes aegypti* but not in *Anopheles maculipennis* mosquitoes following intrathoracic inoculation. Virus did not multiply in experimentally inoculated *Ixodes ricinus* ticks (17).

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**Section X - Histopathology**

Character of lesions (specify host)  
**In man, see references 18,19.**

Inclusion Bodies Intranuclear  
**Man**

Organs/Tissues Affected  
**Brain (M), spinal cord (M), lungs (M,LV), liver (M,LV), spleen (M,LV) kidney (M,LV) heart (M), blood vessels (M, LV), marrow (M), skeletal muscles (M), secretory glands (M)**

Category of tropism

**Section XI - Human Disease**

In Nature Residual Death  
**Significant** **Significant**

Subclinical Overt Disease  
**Significant**

Clinical Manifestations  
**Fever, headache, prostration, conjunctival inflammation, myalgia, arthralgia, CNS signs, hemorrhagic signs, leukopenia, rash, lymphadenopathy, vomiting, hepatitis and diarrhea.**

Number of Cases Category (i.e. febrile illness, etc.)  
**31 (7 deaths)** **Febrile illness with rash, hem. Fever**

**Section XII - Geographic Distribution**

Known (Virus detected)  
**W. Germany , Yugoslavia, and Southern Africa (23)**

Suspected (Antibody only detected)  
**Unknown**

### Section XIII - References

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### Remarks