

#### **Proposed Update of Isolation Precautions Appendix A for Selected High-Consequence Pathogens**

Aaron Kofman, MD

**Prevention and Response Branch** 

**Division of Healthcare Quality Promotion** 

June 8, 2023 – HICPAC Meeting

### Agenda

- 1. Rationale for Update
- 2. Review of VHF and patient placement recommendations (Appendix A)
- 3. Review of IPC considerations for select high-consequence pathogens
  - Marburg, CCHF, Lassa, South American Hemorrhagic Fevers, Andes, Nipah
  - Goal: Inform recommendation update for Appendix A
- 4. Discussion + vote on recommended precautions for each pathogen

## **1.** Rationale for Update

#### **Recent inquiries**

- Marburg outbreaks in Equatorial Guinea, Tanzania (2023)
- 2 U.S. patients with Nipah in differential (2023)
- Lassa, CCHF are often in differential for ill returning travelers from endemic regions
- US had imported Andes virus case (person-to-person transmissible hantavirus) (2018)

 $\rightarrow$  Need for updated recommendations for healthcare infection control precautions

# 2. Review of current VHF PPE and patient placement recommendations

## Appendix A – Viral Hemorrhagic Fevers

Viral hemorrhagic fevers due to Lassa, Ebola, Marburg, Crimean-Congo fever viruses Droplet + Duration of Contact + illness Standard

Ebola Virus Disease for Healthcare Workers [2014] Update: Recommendations for healthcare workers can be found at Ebola For Clinicians. (accessed September 2018).

Single-patient room preferred. Emphasize:

- 1. use of sharps safety devices and safe work practices,
- 2. hand hygiene;
- barrier protection against blood and body fluids upon entry into room (single gloves and fluidresistant or impermeable gown, face/eye protection with masks, goggles or face shields); and

4. appropriate waste handling.

Use N95 or higher respirators when performing aerosolgenerating procedures. Largest viral load in final stages of illness when hemorrhage may occur; additional PPE, including double gloves, leg and shoe coverings may be used, especially in resource-limited settings where options for cleaning and laundry are limited. Notify public health officials immediately if Ebola is suspected [212, 314, 740, 772]. Also see <u>Table 3C</u> for Ebola as a bioterrorism agent.

## **Recommended PPE and Patient Placement for Patients** with suspected EVD (stable or "dry" patients)

While evaluating and managing PUIs who are clinically stable and do not have bleeding, vomiting, or diarrhea, healthcare providers should at a minimum wear:

- Fluid-resistant gown that extends to at least mid-calf or single-use (disposable) fluid-resistant coveralls without integrated hood
- Full face shield
- Facemask
- Gloves with extended cuffs. Two pairs of gloves should be worn. At a minimum, outer gloves should have extended cuffs.

Patient Placement:

- Single patient room with closed door
- AllR for AGPs

# Recommended PPE for unstable/"wet" patients with suspected EVD or confirmed patients with EVD

While evaluating and managing PUIs who are clinically unstable and/or have bleeding, vomiting, or diarrhea, healthcare providers should at a minimum wear:

- Impermeable gown or coverall
- PAPR or N95 respirator
- Examination gloves with extended gloves
- Boot covers (or shoe covers in combination with coverall with integrated socks)
- Apron

**Patient Placement:** 

- Single patient room with closed door
- AllR for AGPs

#### **Proposed Appendix A Update**

- Propose update for the following pathogens
  - VHFs
    - Marburg
    - Lassa
    - Crimean Congo Hemorrhagic Fever (CCHF)
    - South American Arenaviruses (Junin, Machupo, Chapare, Guanarito, Sabia)
    - Andes Virus (a species of hantavirus)
  - Nipah Virus
- Review of virus-specific information to inform decision-making on recommended precautions

3. Review of IPC considerations for select highconsequence pathogens

#### **Appendix A Update - Marburg**

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
	Fever, chills, headache, myalgia, sore	23-90% No vaccine or		Virus has been	Yes	
Marburg	throat, nausea, vomiting May progress to	approved treatments available	Contact with body fluids – blood, most of all	isolated from blood, urine, throat, liver biopsy (autopsy),	Insufficient or no PPE (skin contact with body fluids), sharps injuries,	Same as Ebola
	multi-organ failure, massive hemorrhage	Remdesivir used as treatment, efficacy unclear		eye (anterior chamber)	mucous membrane exposures	

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Appendix A Update – Crimean Congo Hemorrhagic Fever**

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
Crimean Congo Hemorrhagic Fever (CCHF)	Fever, headache, back/joint pain, stomach pain, nausea, vomiting, jaundice Severe bruising, nosebleeds, uncontrolled bleeding at injection sites	3-30% No vaccine or approved treatments available	Contact with body fluids Improper sterilization of medical equipment Percutaneous inoculation from needles Possible droplet/aerosol transmission	PCR detected in blood, nasal swab, saliva, urine, stool, vaginal fluid Viral isolation has been reported from patients/corpses	Yes Percutaneous and cutaneous transmission Possible droplet/aerosol transmission	Same as Ebola

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Appendix A Update – Lassa Fever**

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
	Mild symptoms: flu- like illness	Hospitalized patients' mortality rate: 15-20%	Prolonged contact in setting of unknown exposure			
Lassa	Severe illness: hemorrhage, respiratory distress, vomiting, hearing loss, tremors, encephalitis, multi- organ failure	Overall mortality rate: 1% Ribavirin used as treatment, efficacy unclear No vaccine available	Respiratory droplet or aerosol spread in earlier outbreaks were implicated when source was unknown	Viral culture positive in blood, urine, saliva, and semen	Yes Insufficient or no PPE (skin contact with body fluids)	Same as Ebola

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Appendix A Update – South American Hemorrhagic Fevers**

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally -acquired transmission in healthcare	Proposed PPE and Patient Placement
	All: flu-like illness					
	Junin: absence of respiratory symptoms	Junin: 15-30%, 1% w/ Rx	Junin: P2P transmission surmised in large-scale	Junin: reported from oral swabs, urine, breastmilk, ?sexual		
	Machupo: may develop	Machupo: 25%	outbreaks	transmission	Junin: none	
South American Hemorrhagic Fevers (Arenaviruses)	neurologic/hemorrhagic manifestations	Chapare: 60%	Machupo: P2P transmission demonstrated in 1971, large-	Machupo: blood/throat swab/post-mortem liver/spleen	Machupo: yes	
Junin (Argentine HF)	Chapare: may develop	Guanarito: 33%	scale outbreaks	(viral cx)	Chapare: yes	Same as Ebola
Machupo (Bolivian HF) Chapare (Chapare HF)	ARDS/multiorgan dysfunction	Sabia: 50%	Guanarito: Unclear; only one	Chapare: 2019 outbreak w/		
Guanarito (Venezuelan HF)	Guanarito: respiratory		case of secondary transmission has been identified	blood/urine/conjunctival/semen NP/OP +PCR and culture/NGS	Guanarito: none	
Sabia (Brazilian HF)	symptoms, may develop neurological/hemorrhagic manifestations	Only Junin has vaccine (not available in US)	Chapare: Yes, via contact with body fluids (all)	Guanarito: not established	Sabia: two lab accidents	
	Sabia: may develop multiorgan dysfunction	No proven treatments for any	Sabia: not established	Sabia: not established		

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Appendix A Update – Andes**

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
Andes	Fever, chills, headaches Cough, shortness of breath Respiratory failure Coagulopathy Multiorgan dysfunction	30% No vaccine/treatment	P2P transmission is well-documented among those with close and prolonged contact to case-patients	Breastmilk (PCR) Blood/serum/PBMC (PCR; viral isolate) Urine (PCR) Respiratory sample (PCR)	Yes No/incomplete PPE	Gown, single pair of gloves, respirator, eye protection Patient placement in AIIR

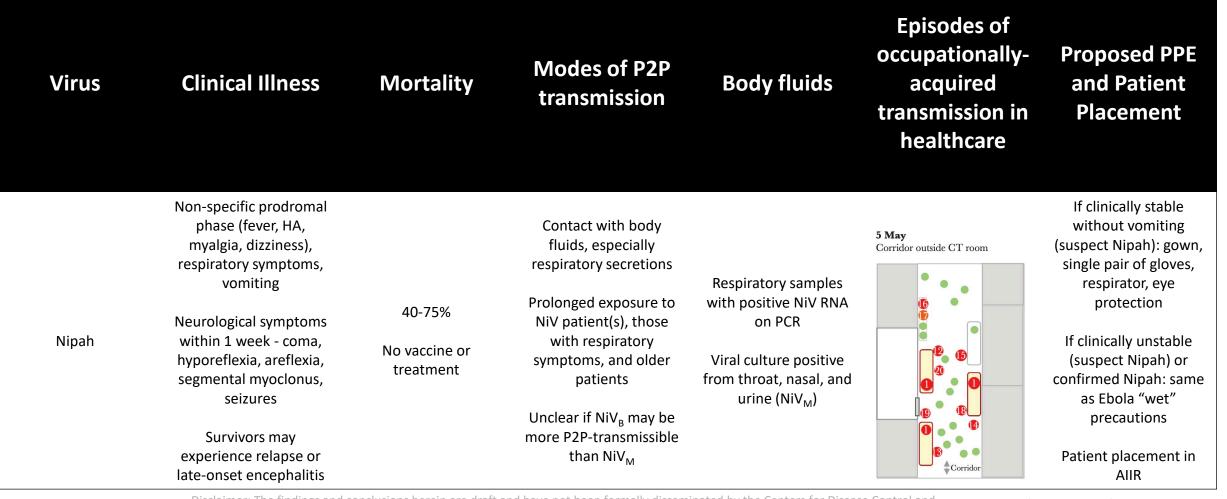
#### **Appendix A Update – Nipah**

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
Nipah	Non-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomiting Neurological symptoms within 1 week - coma,	40-75% No vaccine or	Contact with body fluids, especially respiratory secretions Prolonged exposure to NiV patient(s), those with respiratory	Respiratory samples with positive NiV RNA on PCR	Yes Absence of any or	If clinically stable without vomiting (suspect Nipah): gown, single pair of gloves, respirator, eye protection If clinically unstable
	hyporeflexia, areflexia, segmental myoclonus, seizures Survivors may experience relapse or	treatment	symptoms, and older patients Unclear if NiV <sub>B</sub> may be more P2P-transmissible than NiV <sub>M</sub>	Viral culture positive from throat, nasal, and urine (NiV <sub>M</sub> )	minimal PPE	(suspect Nipah) or confirmed Nipah: same as Ebola "wet" precautions Patient placement in

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and

Prevention and should not be construed to represent any agency determination or policy.

#### Appendix A Update – Nipah (2)



Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

Arunkumar et al JID 2019

## **Discussion + Vote**

#### Appendix A Update – Marburg (2)

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
	Fever, chills, headache, myalgia, sore throat, nausea,	23-90% No vaccine or approved		Virus has been isolated from	Yes Insufficient or no	
Marburg	vomiting May progress to multi-organ failure, massive hemorrhage	treatments available Remdesivir used as treatment, efficacy unclear	Contact with body fluids – blood, most of all	blood, urine, throat, liver biopsy (autopsy), eye (anterior chamber)	PPE (skin contact with body fluids), sharps injuries, mucous membrane exposures	Same as Ebola

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Discussion + Vote on Proposed Update for Marburg**

- Proposal: Change recommended PPE and placement for Marburg to be same as recommended for Ebola
- If change is accepted:
  - Appendix A will be updated to refer to Ebola guidance
  - Ebola guidance will also be updated to include other pathogens to which it applies in addition to Ebola

#### **Appendix A Update – Crimean Congo Hemorrhagic Fever** (2)

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
Crimean Congo Hemorrhagic Fever (CCHF)	Fever, headache, back/joint pain, stomach pain, nausea, vomiting, jaundice Severe bruising, nosebleeds, uncontrolled bleeding at injection sites	3-30% No vaccine or approved treatments available	Contact with body fluids Improper sterilization of medical equipment Percutaneous inoculation from needles Possible droplet/aerosol transmission	PCR detected in blood, nasal swab, saliva, urine, stool, vaginal fluid Viral isolation has been reported from patients/corpses	Yes Percutaneous and cutaneous transmission Possible droplet/aerosol transmission	Same as Ebola

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Discussion + Vote on Proposed Update for CCHF**

- Proposal: Change recommended PPE and placement for CCHF to be same as recommended for Ebola
- If change is accepted:
  - Appendix A will be updated to refer to Ebola guidance
  - Ebola guidance will also be updated to include other pathogens to which it applies in addition to Ebola

#### Appendix A Update – Lassa Fever (2)

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
	Mild symptoms: flu- like illness	Hospitalized patients' mortality rate: 15-20%	Prolonged contact in setting of unknown exposure			
Lassa	Severe illness: hemorrhage, respiratory distress, vomiting, hearing loss, tremors, encephalitis, multi- organ failure	Overall mortality rate: 1% Ribavirin used as treatment, efficacy unclear No vaccine available	Respiratory droplet or aerosol spread in earlier outbreaks were implicated when source was unknown	Viral culture positive in blood, urine, saliva, and semen	Yes Insufficient or no PPE (skin contact with body fluids)	Same as Ebola

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

#### **Discussion + Vote on Proposed Update for Lassa**

- Proposal: Change recommended PPE and placement for Lassa to be same as recommended for Ebola
- If change is accepted:
  - Appendix A will be updated to refer to Ebola guidance
  - Ebola guidance will also be updated to include other pathogens to which it applies in addition to Ebola

#### Appendix A Update – South American Hemorrhagic Fevers (2)

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally -acquired transmission in healthcare	Proposed PPE and Patient Placement
	All: flu-like illness					
	Junin: absence of respiratory symptoms	Junin: 15-30%, 1% w/ Rx	Junin: P2P transmission surmised in large-scale	Junin: reported from oral swabs, urine, breastmilk, ?sexual		
	Machupo: may develop	Machupo: 25%	outbreaks	transmission	Junin: none	
South American Hemorrhagic Fevers (Arenaviruses)	neurologic/hemorrhagic manifestations	Chapare: 60%	Machupo: P2P transmission demonstrated in 1971, large-	Machupo: blood/throat swab/post-mortem liver/spleen	Machupo: yes	
Junin (Argentine HF)	Chapare: may develop	Guanarito: 33%	scale outbreaks	(viral cx)	Chapare: yes	Same as Ebola
Machupo (Bolivian HF) Chapare (Chapare HF) Guanarito (Venezuelan HF)	ARDS/multiorgan dysfunction Guanarito: respiratory	Sabia: 50%	Guanarito: Unclear; only one case of secondary transmission	Chapare: 2019 outbreak w/ blood/urine/conjunctival/semen	Guanarito: none	
Sabia (Brazilian HF)	symptoms, may develop	Only Junin has vaccine (not	has been identified	NP/OP +PCR and culture/NGS	Sabia: two lab accidents	
	neurological/hemorrhagic manifestations	available in US)	Chapare: Yes, via contact with body fluids (all)	Guanarito: not established		
	Sabia: may develop multiorgan dysfunction	No proven treatments for any	Sabia: not established	Sabia: not established		

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and

Prevention and should not be construed to represent any agency determination or policy.

## Discussion + Vote on Proposed Update for South American Hemorrhagic Fevers

- Proposal: Change recommended PPE and placement for South American Hemorrhagic Fevers to be same as recommended for Ebola
- If change is accepted:
  - Appendix A will be updated to refer to Ebola guidance
  - Ebola guidance will also be updated to include other pathogens to which it applies in addition to Ebola

#### Appendix A Update – Andes (2)

Virus	Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	Episodes of occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
Andes	Fever, chills, headaches Cough, shortness of breath Respiratory failure Coagulopathy Multiorgan dysfunction	30% No vaccine/treatment	P2P transmission is well-documented among those with close and prolonged contact to case-patients	Breastmilk (PCR) Blood/serum/PBMC (PCR; viral isolate) Urine (PCR) Respiratory sample (PCR)	Yes No/minimal PPE	Same as Ebola "dry" precautions with the exception of: respirator, single pair of gloves and patient placement in AIIR Patient placement in AIIR

Disclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control and Prevention and should not be construed to represent any agency determination or policy.

### **Discussion + Vote on Proposed Update for Andes**

- Proposal:
  - Same as Ebola "dry" precautions with the exception of: respirator, single pair of gloves and patient placement in AIIR
- If change is accepted:
  - Andes would be added to Appendix A as above

#### Appendix A Update – Nipah (3)

Clinical Illness	Mortality	Modes of P2P transmission	Body fluids	occupationally- acquired transmission in healthcare	Proposed PPE and Patient Placement
Non-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomiting Neurological symptoms	40-75%	Contact with body fluids, especially respiratory secretions Prolonged exposure to NiV patient(s), those	Respiratory samples with positive NiV RNA on PCR	Yes	If clinically stable without vomiting (suspect Nipah): same as Ebola "dry" precautions with exception of single pair of gloves, respirator, and patient placement
within 1 week - coma, hyporeflexia, areflexia, segmental myoclonus, seizures Survivors may experience relapse or late-onset encephalitis	No vaccine or treatment	with respiratory symptoms, and older patients Unclear if NiV <sub>B</sub> may be more P2P-transmissible than NiV <sub>M</sub>	Viral culture positive from throat, nasal, and urine (NiV <sub>M</sub> )	Absence of any or minimal PPE	in AIIR If clinically unstable or confirmed Nipah: same as Ebola "wet" precautions and patient placement in AIIR
	Non-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomiting Neurological symptoms within 1 week - coma, hyporeflexia, areflexia, segmental myoclonus, seizures Survivors may experience relapse or late-onset encephalitis Disclaimer: The findings and c	Non-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomiting Neurological symptoms within 1 week - coma, hyporeflexia, areflexia, segmental myoclonus, seizures Survivors may experience relapse or late-onset encephalitis Disclaimer: The findings and conclusions herein are draft	Clinical IllnessMortalitytransmissionNon-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomitingContact with body fluids, especially respiratory secretionsNeurological symptoms within 1 week - coma, hyporeflexia, areflexia, segmental myoclonus, seizures40-75%Neurological symptoms within 1 week - coma, hyporeflexia, areflexia, seizures40-75%No vaccine or treatmentProlonged exposure to NiV patient(s), those with respiratory symptoms, and older patientsSurvivors may experience relapse or late-onset encephalitisUnclear if NiV <sub>B</sub> may be more P2P-transmissible than NiV <sub>M</sub>	Clinical IllnessMortalityBody fluidsNon-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomitingContact with body fluids, especially respiratory secretionsRespiratory samples with n 1 week - coma, hyporeflexia, areflexia, seizuresNeurological symptoms, within 1 week - coma, hyporeflexia, areflexia, seizures40-75%Contact with body fluids, especially respiratory secretionsRespiratory samples with positive NiV RNA on PCRNo vaccine or treatmentNo vaccine or treatmentViral culture positive from throat, nasal, and urine (NiV_M)Survivors may experience relapse or late-onset encephalitisSurvivors may than NiV_MUnclear if NiV <sub>B</sub> may be more P2P-transmissible than NiV_M	Clinical IllnessMortalityModes of P2P transmissionBody fluidsacquired transmission in healthcareNon-specific prodromal phase (fever, HA, myalgia, dizziness), respiratory symptoms, vomitingContact with body fluids, especially respiratory secretionsRespiratory samples with positive NiV RNA on PCRYesNeurological symptoms within 1 week - coma, hyporeflexia, areflexia, segurental myoclonus, seizures40-75% No vaccine or treatmentProlonged exposure to NiV patient(s), those with respiratory symptoms, and older patientsRespiratory samples with positive NiV RNA on PCRYesSurvivors may experience relapse or late-onset encephalitisUnclear if NiV <sub>B</sub> may be more P2P-transmissible than NiV <sub>M</sub> Viral culture positive from throat, nasal, and urine (NiV <sub>M</sub> )Absence of any or minimal PPEDisclaimer: The findings and conclusions herein are draft and have not been formally disseminated by the Centers for Disease Control andMortal table

Prevention and should not be construed to represent any agency determination or policy.

## **Discussion + Vote on Proposed Update for Nipah**

- Proposal
  - If clinically stable without vomiting (suspect Nipah): same as Ebola "dry" precautions with exception of single pair of gloves, respirator, and patient placement in AIIR
  - If clinically unstable or confirmed Nipah: same as Ebola "wet" precautions and patient placement in AIIR
- If change is accepted:
  - Nipah would be added to Appendix A as above

# Thank you!

#### Acknowledgments

- David Kuhar
- Melissa Schaefer
- Alex Kallen
- Ryan Fagan
- Joe Perz
- Caitlin Cossaboom



#### **Key references**

#### Marburg

- Bausch DG, Borchert M, Grein T, Roth C, Swanepoel R, Libande ML, Talarmin A, Bertherat E, Muyembe-Tamfum JJ, Tugume B, Colebunders R. Risk factors for Marburg hemorrhagic fever, Democratic Republic of the Congo. Emerging Infectious Diseases. 2003 Dec;9(12):1531.
- Gear JS, Cassel GA, Gear AJ, Trappler B, Clausen L, Meyers AM, Kew MC, Bothwell TH, Sher R, Miller GB, Schneider J. Outbreak of Marburg virus disease in Johannesburg. Br Med J. 1975 Nov 29;4(5995):489-93.
- Martini GA. Marburg virus disease. Postgraduate medical journal. 1973 Aug 1;49(574):542-6.
- Brainard J, Pond K, Hooper L, Edmunds K, Hunter P. Presence and persistence of Ebola or Marburg virus in patients and survivors: a rapid systematic review. PLoS neglected tropical diseases. 2016 Feb 29;10(2):e0004475.
- Selvaraj SA, Lee KE, Harrell M, Ivanov I, Allegranzi B. Infection rates and risk factors for infection among health workers during Ebola and Marburg virus outbreaks: a systematic review. The Journal of infectious diseases. 2018 Nov 22;218(suppl\_5):S679-89.
- Smith DH, Isaacson M, Johnson KM, Bagshawe A, Johnson BK, Swanapoel R, Killey M, Siongok T, Keruga WK. Marburg-virus disease in Kenya. The Lancet. 1982 Apr 10;319(8276):816-20.

## **Key references CCHF**

#### CCHF

- Gozel MG, Dokmetas I, Oztop AY, Engin A, Elaldi N, Bakir M. Recommended precaution procedures protect healthcare workers from Crimean-Congo hemorrhagic fever virus. International Journal of Infectious Diseases. 2013 Nov 1;17(11):e1046-50.
- Celikbas AK, Dokuzoğuz B, Baykam N, Gok SE, Eroğlu MN, Midilli K, Zeller H, Ergonul O. Crimean-Congo hemorrhagic fever among health care workers, Turkey. Emerging infectious diseases. 2014 Mar;20(3):477.
- Altaf A, Luby S, Jamil A, Najam A, Aamir Z, Khan J, Mirza S, McCormick J, Fisher-Hoch S. Outbreak of Crimean-Congo haemorrhagic fever in Quetta, Pakistan: contact tracing and risk assessment. Tropical Medicine & International Health. 1998 Nov;3(11):878-82.
- Maltezou HC, Maltezos E, Papa A. Contact tracing and serosurvey among healthcare workers exposed to Crimean-Congo haemorrhagic fever in Greece. Scandinavian journal of infectious diseases. 2009 Jan 1;41(11-12):877-80.
- Leblebicioglu H, Sunbul M, Guner R, Bodur H, Bulut C, Duygu F, Elaldi NA, Senturk GC, Ozkurt Z, Yilmaz G, Fletcher TE. Healthcare-associated Crimean-Congo haemorrhagic fever in Turkey, 2002–2014: a multicentre retrospective cross-sectional study. Clinical microbiology and infection. 2016 Apr 1;22(4):387-e1.
- Pshenichnaya NY, Nenadskaya SA. Probable Crimean-Congo hemorrhagic fever virus transmission occurred after aerosol-generating medical procedures in Russia: nosocomial cluster. International Journal of Infectious Diseases. 2015 Apr 1;33:120-2.
- Tsergouli K, Karampatakis T, Haidich AB, Metallidis S, Papa A. Nosocomial infections caused by Crimean–Congo haemorrhagic fever virus. Journal of Hospital Infection. 2020 May 1;105(1):43-52.
- Yildirmak T, Tulek N, Bulut C. Crimean–Congo haemorrhagic fever: transmission to visitors and healthcare workers. Infection. 2016 Oct;44:687-9.
- Yagci-Caglayik D, Kayaaslan B, Yapar D, Kocagul-Celikbas A, Ozkaya-Parlakay A, Emek M, Baykam N, Tezer H, Korukluoglu G, Ozkul A. Monitoring Crimean-Congo haemorrhagic fever virus RNA shedding in body secretions and serological status in hospitalised patients, Turkey, 2015. Eurosurveillance. 2020 Mar 12;25(10):1900284.
- Bodur H, Akıncı E, Öngürü P, Carhan A, Uyar Y, Tanrıcı A, Cataloluk O, Kubar A. Detection of Crimean-Congo hemorrhagic fever virus genome in saliva and urine. International Journal of Infectious Diseases. 2010 Mar 1;14(3):e247-9.
- Hoogstraal H. The epidemiology of tick-borne Crimean-Congo hemorrhagic fever in Asia, Europe, and Africa. Journal of medical entomology. 1979 May 22;15(4):307-417.
- Burt FJ, Swanepoel R, Shieh WJ, Smith JF. Immunohistochemical and in situ localization of Crimean-Congo hemorrhagic fever (CCHF) virus in human tissues and implications for CCHF pathogenesis. Archives of pathology & laboratory medicine. 1997 Aug 1;121(8):839.

#### **Key references Lassa**

#### Lassa

- Carey DE, Kemp GE, White HA, Pinneo L, Addy RF, Fom AL, Stroh G, Casals J, Henderson BE. Lassa fever epidemiological aspects of the 1970 epidemic, Jos, Nigeria. Transactions of the Royal Society of Tropical Medicine and Hygiene. 1972 Jan 1;66(3):402-8.
- Monath TP, Casals J. Diagnosis of Lassa fever and the isolation and management of patients. Bulletin of the World Health Organization. 1975;52(4-6):707.
- Monath TP. A hospital epidemic of Lassa fever, in Zorzor, Liberia, March-April 1972. American journal of tropical medicine and hygiene. 1973;22(6):773-9.
- Ajayi NA, Nwigwe CG, Azuogu BN, Onyire BN, Nwonwu EU, Ogbonnaya LU, Onwe FI, Ekaete T, Günther S, Ukwaja KN. Containing a Lassa fever epidemic in a resource-limited setting: outbreak description and lessons learned from Abakaliki, Nigeria (January–March 2012). International Journal of Infectious Diseases. 2013 Nov 1;17(11):e1011-6.
- Ehlkes L, George M, Samosny G, Burckhardt F, Vogt M, Bent S, Jahn K, Zanger P. Management of a lassa fever outbreak, Rhineland-Palatinate, Germany, 2016. Eurosurveillance. 2017 Sep 28;22(39):16-00728.
- Haas WH, Breuer T, Pfaff G, Schmitz H, Köhler P, Asper M, Emmerich P, Drosten C, Gölnitz U, Fleischer K, Günther S. Imported Lassa fever in Germany: surveillance and management of contact persons. Clinical infectious diseases. 2003 May 15;36(10):1254-8.
- Dan-Nwafor CC, Ipadeola O, Smout E, Ilori E, Adeyemo A, Umeokonkwo C, Nwidi D, Nwachukwu W, Ukponu W, Omabe E, Anaebonam U. A cluster of nosocomial Lassa fever cases in a tertiary health facility in Nigeria: Description and lessons learned, 2018. International Journal of Infectious Diseases. 2019 Jun 1;83:88-94.
- Fisher-Hoch SP, Tomori O, Nasidi A, Perez-Oronoz GI, Fakile Y, Hutwagner L, McCormick JB. Review of cases of nosocomial Lassa fever in Nigeria: the high price of poor medical practice. Bmj. 1995 Sep 30;311(7009):857-9.
- Fisher-Hoch SP, Craven RB, Forthall DN, Scott SM, Price ME, Price FM, Sasso DR, McCormick JB. Safe intensive-care management of a severe case of Lassa fever with simple barrier nursing techniques. The Lancet. 1985 Nov 30;326(8466):1227-9.
- Grahn A, Bråve A, Tolfvenstam T, Studahl M. Absence of nosocomial transmission of imported Lassa fever during use of standard barrier nursing methods. Emerging Infectious Diseases. 2018 Jun;24(6):972.
- Helmick C, Scribner C, Webb P, Krebs J, Mccormick J. No evidence for increased risk of Lassa fever infection in hospital staff. The Lancet. 1986 Nov 22;328(8517):1202-5.
- Raabe V, Koehler J. Laboratory diagnosis of Lassa fever. Journal of Clinical Microbiology. 2017 Jun;55(6):1629-37.

#### **Key references SAHF**

#### **South American Hemorrhagic Fevers**

- Frank MG, Beitscher A, Webb CM, Raabe V, Bhadelia N, Cieslak TJ, Davey RT, Dierberg K, Evans JD, Grein J, Kortepeter MG. South American hemorrhagic fevers: a summary for clinicians. International Journal of Infectious Diseases. 2021 Apr 1;105:505-15.
- Barry M, Russi M, Armstrong L, Geller D, Tesh R, Dembry L, Gonzalez JP, Khan AS, Peters CJ. Treatment of a laboratory-acquired Sabia virus infection. New England Journal of Medicine. 1995 Aug 3;333(5):294-6.
- Armstrong LR, Dembry LM, Rainey PM, Russi MB, Khan AS, Fischer SH, Edberg SC, Ksiazek TG, Rollin PE, Peters CJ. Management of a Sabia virus-infected patient in a US hospital. Infection Control & Hospital Epidemiology. 1999 Mar;20(3):176-82.
- Manzione ND, Salas RA, Paredes H, Godoy O, Rojas L, Araoz F, Fulhorst CF, Ksiazek TG, Mills JN, Ellis BA, Peters CJ. Venezuelan hemorrhagic fever: clinical and epidemiological studies of 165 cases. Clinical infectious diseases. 1998 Feb 1;26(2):308-13.
- Peters CJ, Kuehne RW, Mercado RR, Le Bow RH, Spertzel RO, Webb PA. Hemorrhagic fever in cochabamba, bolivia, 1971. American journal of epidemiology. 1974 Jun 1;99(6):425-33.
- Salas R, Pacheco ME, Ramos B, Taibo ME, Jaimes E, Vasquez C, Querales J, de Manzione N, Godoy O, Betancourt A, Araoz F. Venezuelan haemorrhagic fever. The Lancet. 1991 Oct 26;338(8774):1033-6.
- Veliziotis I, Roman A, Martiny D, Schuldt G, Claus M, Dauby N, Van den Wijngaert S, Martin C, Nasreddine R, Perandones C, Mahieu R. Clinical management of Argentine hemorrhagic fever using ribavirin and favipiravir, Belgium, 2020. Emerging infectious diseases. 2020 Jul;26(7):1562.
- Loayza Mafayle R, Morales-Betoulle ME, Romero C, Cossaboom CM, Whitmer S, Alvarez Aguilera CE, Avila Ardaya C, Cruz Zambrana M, Dávalos Anajia A, Mendoza Loayza N, Montaño AM. Chapare hemorrhagic fever and virus detection in rodents in Bolivia in 2019. New England Journal of Medicine. 2022 Jun 16;386(24):2283-94.
- Ellwanger JH, Chies JA. Keeping track of hidden dangers-The short history of the Sabiá virus. Revista da Sociedade Brasileira de Medicina Tropical. 2017 Jan;50:03-8.
- Enria DA, Briggiler AM, Sánchez Z. Treatment of Argentine hemorrhagic fever. Antiviral research. 2008 Apr 1;78(1):132-9.

## Key references Nipah

#### Nipah

- Arunkumar G, Chandni R, Mourya DT, Singh SK, Sadanandan R, Sudan P, Bhargava B. Outbreak investigation of Nipah virus disease in Kerala, India, 2018. The Journal of infectious diseases. 2019 May 24;219(12):1867-78.
- Chadha MS, Comer JA, Lowe L, Rota PA, Rollin PE, Bellini WJ, Ksiazek TG, Mishra AC. Nipah virus-associated encephalitis outbreak, Siliguri, India. Emerging infectious diseases. 2006 Feb;12(2):235.
- Gurley ES, Montgomery JM, Hossain MJ, Bell M, Azad AK, Islam MR, Molla MA, Carroll DS, Ksiazek TG, Rota PA, Lowe L. Person-toperson transmission of Nipah virus in a Bangladeshi community. Emerging infectious diseases. 2007 Jul;13(7):1031.
- Gurley ES, Montgomery JM, Hossain MJ, Islam MR, Molla MA, Shamsuzzaman SM, Akram K, Zaman K, Asgari N, Comer JA, Azad AK. Risk of nosocomial transmission of Nipah virus in a Bangladesh hospital. Infection Control & Hospital Epidemiology. 2007 Jun;28(6):740-2.
- Hegde S, Lee KH, Styczynski A, Jones F, Gomes I, Das P, Gurley ES. Potential for person-to-person transmission of henipaviruses: A systematic review of the literature. medRxiv. 2023:2023-02.
- Hughes JM, Wilson ME, Luby SP, Gurley ES, Hossain MJ. Transmission of human infection with Nipah virus. Clinical Infectious Diseases. 2009 Dec 1;49(11):1743-8.
- Chua KB, Lam SK, Goh KJ, Hooi PS, Ksiazek TG, Kamarulzaman A, Olson J, Tan CT. The presence of Nipah virus in respiratory secretions and urine of patients during an outbreak of Nipah virus encephalitis in Malaysia. Journal of Infection. 2001 Jan 1;42(1):40-3.
- Nikolay B, Salje H, Hossain MJ, Khan AD, Sazzad HM, Rahman M, Daszak P, Ströher U, Pulliam JR, Kilpatrick AM, Nichol ST. Transmission of Nipah virus—14 years of investigations in Bangladesh. New England Journal of Medicine. 2019 May 9;380(19):1804-14.

#### **Key references Andes**

#### Andes

- Martínez VP, Di Paola N, Alonso DO, Pérez-Sautu U, Bellomo CM, Iglesias AA, Coelho RM, López B, Periolo N, Larson PA, Nagle ER. "Super-spreaders" and person-to-person transmission of Andes virus in Argentina. New England Journal of Medicine. 2020 Dec 3;383(23):2230-41.. 2019 May 9;380(19):1804-14.
- Chaparro J, Vega J, Terry W, Vera JL, Barra B, Meyer R, Peters CJ, Khan AS, Ksiazek TG. Assessment of person-to-person transmission of hantavirus pulmonary syndrome in a Chilean hospital setting. Journal of Hospital Infection. 1998 Dec 1;40(4):281-5.
- Ferrés M, Vial P, Marco C, Yanez L, Godoy P, Castillo C, Hjelle B, Delgado I, Lee SJ, Mertz GJ, Andes Virus Household Contacts Study Group. Prospective evaluation of household contacts of persons with hantavirus cardiopulmonary syndrome in Chile. The Journal of infectious diseases. 2007 Jun 1;195(11):1563-71.
- Kofman A, Eggers P, Kjemtrup A, Hall R, Brown SM, Morales-Betoulle M, Graziano J, Zufan SE, Whitmer SL, Cannon DL, Chiang CF. Notes from the field: contact tracing investigation after first case of andes virus in the United States—Delaware, February 2018. Morbidity and Mortality Weekly Report. 2018 Oct 10;67(41):1162.
- Kuenzli AB, Marschall J, Schefold JC, Schafer M, Engler OB, Ackermann-Gäumann R, Reineke DC, Suter-Riniker F, Staehelin C. Hantavirus cardiopulmonary syndrome due to imported Andes hantavirus infection in Switzerland: a multidisciplinary challenge, two cases and a literature review. Clinical infectious diseases. 2018 Nov 13;67(11):1788-95.
- Martinez VP, Bellomo C, San Juan J, Pinna D, Forlenza R, Elder M, Padula PJ. Person-to-person transmission of Andes virus. Emerging infectious diseases. 2005 Dec;11(12):1848.
- Martinez-Valdebenito C, Calvo M, Vial C, Mansilla R, Marco C, Palma RE, Vial PA, Valdivieso F, Mertz G, Ferrés M. Person-to-person household and nosocomial transmission of Andes hantavirus, Southern Chile, 2011. Emerging infectious diseases. 2014 Oct;20(10):1629.
- Padula PJ, Edelstein A, Miguel SD, Lopez NM, Rossi CM, Rabinovich RD. Hantavirus pulmonary syndrome outbreak in Argentina: molecular evidence for person-to-person transmission of Andes virus. Virology. 1998 Feb 15;241(2):323-30.
- Toledo J, Haby MM, Reveiz L, Sosa Leon L, Angerami R, Aldighieri S. Evidence for human-to-human transmission of hantavirus: a systematic review. The Journal of infectious diseases. 2022 Oct 15;226(8):1362-71.