



Monkeypox Virus: A Review

[Huda Mawlood Taher*](#), [Hiyam Jamal Ibrahim](#), [Bayan Mohammed Mahdi](#)

Department of Biology, College of Education for Pure Sciences, University of Kirkuk, Iraq.

*Corresponding Author: huda.mawlood@uokirkuk.edu.iq

Citation: Taher HM, Ibrahim HJ, Mahdi BM. Monkeypox Virus: A Review. Al-Kitab J. Pure Sci. [Internet]. 2025 May. 02 ;9(2):128-139. DOI: <https://doi.org/10.32441/kjps.09.02.p8>.

Keywords: Monkeypox, virus, Pathogenicity.

Article History

Received	14 Oct. 2024
Accepted	04 Dec. 2024
Available online	02 May. 2025

©2025. THIS IS AN OPEN-ACCESS ARTICLE UNDER THE CC BY LICENSE
<http://creativecommons.org/licenses/by/4.0/>



Abstract:

Monkeypox is a disease caused by the monkeypox virus, which is related to the smallpox virus. It is a double-strand DNA virus enclosed in an envelope and a member of the Orthopoxvirus genus within the Orthopoxviridae family. This disease was spread in West and Central African countries and other regions as well. It transmits through direct contact with the skin lesions of an infected person, as well as through respiratory droplets and contaminated materials. Monkeypox is generally less severe than smallpox, and most people recover completely from it; however, it is still serious, especially for young children, pregnant women, and people with weakened immune systems. It causes pain, fever, rash, and swollen lymph nodes.

Keywords: Monkeypox, Virus, Pathogenicity.

فيروس جدري القروود: مراجعة

هدى مولود طاهر*، هيام جمال إبراهيم، بيان محمد مهدي

قسم علوم الحياة، كلية التربية للعلوم الصرفة، جامعة كركوك، العراق
huda.mawlood@uokirkuk.edu.iq, plantanatomy@uokirkuk.edu.iq, bavanmohammad@uokirkuk.edu.iq

الخلاصة:

جدري القروود هو مرض يسببه فيروس جدري القروود وهو قريب من فيروس الجدري، وهو فيروس ثنائي السلسلة محاط بغلاف وهو من جنس فيروسات الأورثوبوكس ضمن عائلة الفيروسات الأورثوبوكس. انتشر هذا المرض في دول غرب ووسط أفريقيا ومناطق أخرى أيضًا. ينتقل من خلال الاتصال المباشر بأفات الجلد لدى الشخص المصاب، وكذلك من خلال قطرات الجهاز التنفسي والمواد الملوثة. جدري القروود أقل خطورة من الجدري بشكل عام ويتعافى منه معظم الناس تمامًا، ولكنه لا يزال خطيرًا، وخاصة الأطفال الصغار والنساء الحوامل والأشخاص الذين يعانون من ضعف في جهاز المناعة. حيث يسبب ألمًا وحمى وطفح جلدي وتضخم الغدد اللمفاوية.

الكلمات المفتاحية: جدري القروود، فيروس، أمراضية.

1. Introduction

Monkeypox (MPXV) is a viral infection transmitted from animals to humans that leads to a skin rash resembling smallpox, and the likelihood of death is much lower in monkeypox compared to smallpox [1]. Two chief groups of monkeypox virus have been discovered in West and Central Africa, with the former linked to another serious disease [2]. Numerous instances in the recent epidemic are linked to sexual transmission, particularly between individuals who are recognized as bisexual, gay or engage in sex with men [3]. Firstly, the monkeypox virus was discovered in monkeys [4]. Monkeypox was reported in 1958 in monkeys that were transported from Singapore to Denmark [5]. The initial incident in humans was identified in a 9-month-old boy, Zaire, in 1970 (currently known as the Democratic Republic of the Congo) [6]. Since that period, monkeypox has established itself as a common disease in the DRC where it has expanded to various African nations, particularly in West and Central Africa. The initial documented monkeypox condition outside of Africa was reported in 2003. (97.962)cases of monkeypox have been recorded on June 30, 2024, in nearly 120 countries, 110 of which have not historically announced it [7].

The monkeypox virus comes in two types: clade I and clade II [8]:

- Clade I causes more severe illnesses and deaths. Up to 10% of cases in some outbreaks have resulted in deaths, however, the death toll from more recent outbreaks has been lower. Clade I is prevalent in Central Africa.

- Clade II is the type that caused the global epidemic that began in 2022. It's not as serious. Over 99.9% of people make it out alive. Clade II is prevalent in West Africa.

2. Transmission methods of monkeypox virus

There is currently no known method of MPXV transmission to humans. It is assumed that handling monkeypox-infected animals directly by touch, bite, scratch, and indirectly causes the primary animal-to-human infection [9]. It is believed that the respiratory system, mucous membranes, and damaged skin let the virus enter the body (eyes, nose, and mouth), Secondary human-to-human transmission frequently occurs as a result of respiratory droplets, direct or indirect contact with bodily fluids, solid lesions, contaminated surfaces, and contaminated clothing. Extended interactions with patients increase the risk of infection for hospital staff and family members [10]. Generally, asleep in the same bed or room, alive in an identical home, and eating or drinking from the same dish were hazardous behaviors connected with person-to-person transmission [11]. Also, it has been established that sleeping outside or on the ground and residing close to or visiting a forest increase the likelihood of coming into contact with animals and, consequently, the risk of the monkeypox virus dispersal from animal to human [12]. Using toilet and sterility and washing wear did not have a significant connotation with obtaining monkeypox [13]. During the 2003 occurrence in the US, daily contact with sick animals and cleaning their cages were found to be dangerous factors for developing monkeypox [14].

3. Epidemiology

Human monkeypox infections have been reported in African countries, the most affected countries are Nigeria, the Democratic Republic of the Congo, and the Central African Republic, which has only a few hundred cases reported [15], together with a very great rise in case numbers above the previous thirty years [16], with national observation data showed in assumed cases from 2001 to 2018: from fewer than 500 cases to more than 2500 cases [15]. The majority of cases reported in the 1970s and 1980s involved young children, while the average age at identification in Nigeria during the 2017–2018 epidemic was 20-29 years old [17].

In Africa, both person-to-person and animal-to-person transmissions have been acknowledged. Transmission of zoonotic organisms happens by connection with the lesions or biological fluids of an infested animal [18]. These associates happen during hunting, butchering, or game consumption. Person-to-person transmission occurs mainly by contact with biological fluids, infected skin lesions of patients, and polluted resources such as bedding also can be transferable. The cause of transmission in Africa is unknown, and studies have found

significant differences in the amounts of suspected person-to-person and animal-to-person transmissions. Of the 122 patients in Nigeria, 10 had contact with animals and 36 had contact with people who had an interrelated skin lesion [17]. Intra-familial communications have been reported for viral clades 1 and 2 [19]. Sexual transmission has been reported in uncommon cases in African countries [20]. Chains of transmission are commonly stumpy up to seven transmission goings on within a lone household have been reported [21].

4. Pathogenicity

Symptoms of infection of monkeypox include headaches, body aches, chills, sore throat, fever, fatigue, swollen lymph nodes, malaise, and a severe rash that develops into blisters or papules that crust over and heal [22]. Monkeypox Lymphadenopathy appears in MPXV infections but may be misdiagnosed as chickenpox [23]. Numerous patients experience different diseases including few or no lesions, which are restricted to the genital perianal area, hurt, and hemorrhage [24].

The beginning monkeypox signs varieties start from 6 to 21 days, and naturally, the infection restores to its personnel in 2–4 weeks [25]. Monkeypox Virus infections may cause lengthy viral DNA remaining in the higher respiratory tract that persists later in the skin lesion resolves, but it's inexact if this is connected with communicable virus transmission [26]. The common monkeypox Virus infection patches up on their particular. Though, it is more severe, and potency needs hospitalization in immunosuppressed patients and young children [27]. HIV-1-infected people have lengthy monkeypox disease, greater lesions, secondary microbial skin infections, and genital abscesses [22]. Also, monkeypox Virus may be diffused through the placenta in the pregnancy and clue to fetal death [28].

In several cases, possibly life-threatening problems such as encephalitis secondary infection of the instrument, and bronchopneumonia [23]. An additional rare but serious, long-term problem of monkeypox is the loss of vision resulting from infection of the tissue damage and eye cornea [28]. The overall mortality rate differs depending on patient age, infection types, and localization of the occurrence. Also, it is more fatal in children than in adults [29].

5. Immune response

The first line of defense against a monkeypox infection is thought to be the innate immune system. Innate immune cells, which produce inflammatory cytokines, and type I interferons, such as monocytes and NK cells, play a vital role. However, NK cells and cytotoxicity can be overcome by monkeypox [30]. Th2-related cytokines (IL-4, IL-5, IL-6, and IL-10) are elevated in response to monkeypox infection, whereas Th1-associated cytokines (IL-2, IL-12, TNF- α ,

and IFN- γ) stay within the normal range, suggesting complex immune deregulations during the infection [31]. Antibody response, which is a key component of adaptive immunity, is essential for eliminating monkeypox. Detection of specific IgG and IgM antibodies in infected individuals is commonly used for diagnosis, while B cells play a crucial role in providing protection against monkeypox through vaccination [30]. Also, T cells contribute to the protecting responses to monkeypox, as T cell response to monkeypox has been identified in healthy persons, especially persons born before the year 1976 who likely expected the smallpox vaccine, which affords cross defense against monkeypox [32].

Some factors affect the immune response to monkeypox disease, people who have had smallpox in the past tend to be more immune to monkeypox because of the relationship between the viruses and the presence of cross-immunity, leading to a more robust genetic makeup and immune reaction. Younger people tend to have a stronger immune response to the monkeypox virus compared to older people, due to the immune system declining with age [33]. But immune-compromised persons like persons with cancer, HIV, and organ transplant receivers are at a greater hazard of contracting monkeypox and my knowledge of elongated recovery periods and their conceded immune status hinders their bodies' capability to effectually conflict with the monkeypox virus, as **Figure 1** [34].

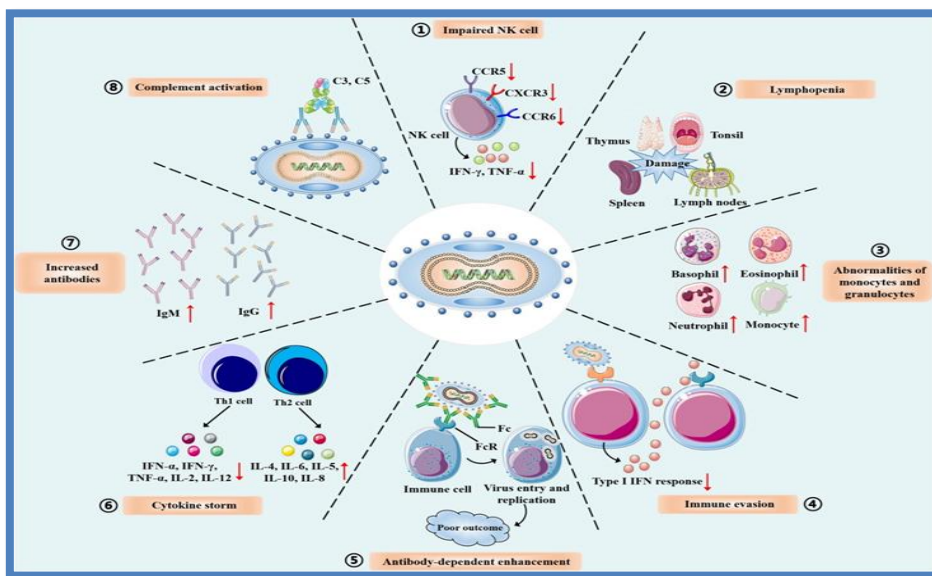


Figure 1: The immunopathogenesis of monkeypox virus [34]

6. Diagnosis methods

Diagnosis is considered the key to detection and keeping monkeypox under control. We can simply distinguish monkeypox from other orthopox viruses by electron microscopy. Clinical characteristics can help differentiate poxvirus infections from other causes of vesiculopustular

eruptions, a laboratory examination is necessary for a conclusive diagnosis [35]. IgM and IgG ELISA, immune fluorescent antibody assay, PCR, electron microscopy, virus isolation, and histopathology analysis are laboratory diagnostic methods for monkeypox [36]. We can distinguish between a herpes virus and poxvirus infection by using immunohistochemistry analysis with monoclonal or polyclonal antibodies against all orthopox viruses [37]. Electron microscopes have played a chief role in viral diagnosis, they can be the main mode for diagnosis of poxvirus infection. We can be detected under electron microscopy the characteristic morphology of poxvirus [36]. The definitive identification of MPXV often involves virus isolation in the culture of mammalian cells and their characterization using different PCR techniques, either by sequencing of amplicons or by restriction fragment length polymorphism analysis [38]. Also, the accessibility of many real-time PCR assays that use monkeypox Virus definite targets has improved in recent years [39]. A new rapid method for identifying orthopox viruses was created using a DNA microarray with the TNF receptor gene crmB [40].

The main diagnostic problem is distinguishing between monkeypox and varicella. Laboratory authorization can be done by PCR technique of vesicle liquid or scab in active disease, however when disease determination, testing of convalescent stage serum specimen for anti varicella virus IgM can be done[36]. Identifying anti-poxvirus antibodies in a person who has not ever been vaccinated and has a past of serious infection and skin rash suggests a diagnosis of monkeypox [40].

7. Prevention

The prevention of the monkeypox virus in healthcare clinicians and specialists is an additional challenge since they are normally in close contact with infected people [41]. To decrease the infection risk, healthcare professionals should follow the commendations such as carefully handled connection with special protective equipment such as gloves, a gown, eye protection, and a fitted N95 mask (**Figure 2**) [42]. Patients with disbelieved monkeypox infections should also be hidden, and their skin lesions must be covered with a gown or cloth, and be placed in isolation, if at all possible, in a single-person room [43]. Patients should remain in the home and border contact with others for 3–6 weeks and evade close contact such as sexual connection with someone exposed or infected with the monkeypox virus, persons should maintain good hand sanitization and respiratory system by wearing a fitted mask and lidding coughs and sneeze with the bend of a limb after taking visitors at home, proper cleaning are advised [44]. The obtainable documents show that smallpox vaccination might ensure 85% cross-protection against monkeypox [45].

Samples such as scabs or cutaneous tissues should be handled with care because the simplicity of transmission by direct contact or air particles, and instantaneous ring vaccination and quarantine are considered the simply actual public health protecting procedures as there is no effective, approved antiviral healing for monkeypox [46].

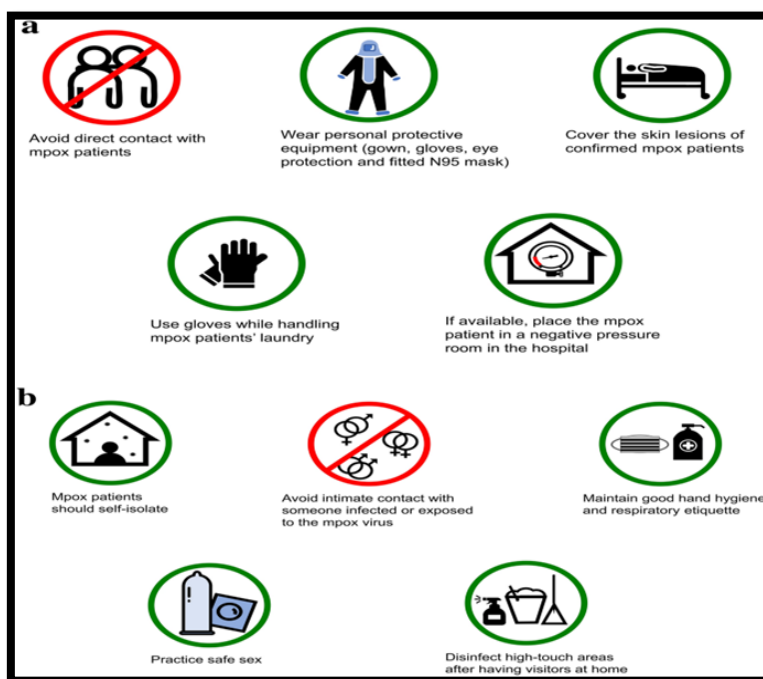


Figure 2: Prevention ways from monkeypox virus [42].

8. Treatment

Most individuals infected with monkeypox infection recover without medicinal treatment, while patients with gastrointestinal warning signs such as nausea, and diarrhea will need oral or venous rehydration to decrease losses of gastrointestinal fluid [47]. The number of antivirals might be effective in handling monkeypox infection [48]. Tecovirimat also known as ST-246 or TPOXX completely that chemical considered the best antiviral directed for treating smallpox in adults and children weighing over 3 kg [49]. Double therapy with brincidofovir and tecovirimat can be used in patients with severe infection. Tecovirimat mechanisms inhibit viral envelope protein VP37, which controls the final steps of virus maturation and release from infected cells, thereby inhibiting virus spread within the infected host [50]. Even though the efficiency of this agent in persons contrary to monkeypox has not been tried, revisions have informed advanced persistence from deadly monkeypox virus infections in recoverin at treated animals associated to palliative treated animals at changed periods of disease [51]. In some research studies, tecobilimatosi has been used in combination with vaccinia immune globulin in patients with difficult-to-administer smallpox vaccination, such as those with vaccinia

eczema and advanced vaccine reactions [52]. The CDC New protocol enables us of Tecovirimat in non-variola orthopoxvirus infections as monkeypox [53]. It also includes information for pediatric patients under 13 kg to open an oral capsule and combine the liquid or soft food inside of it [54]. It was also approved to utilize brincidofovir for treating smallpox in the US in the meantime in 2021 [55]. Brincidofovir by oral considered an equivalent to the intravenous drug cidofovir, a lesser amount of kidney toxicity related to cidofovir, these medicines work through inhibit the viral DNA polymerase [56]. Clinical documents about the efficacy of cidofovir against monkeypox in humans are unknown, but the in vitro activity and efficacy of cidofovir against lethal monkeypox virus infections in animals are known. The efficacy of cidofovir against monkeypox in humans is missing, yet in vitro activity and efficiency against fatal monkeypox virus infections in animals have been informed [57]. Liver function examinations must be prepared before and through treatment with brincidofovir, these medications may lead to a rise the serum transaminases and serum bilirubin. It can use Vaccinia hyperimmune globulin which is qualified by the FDA for cooperation of certain complications of vaccination [58]. VIG subsidy for patients with severe immunodeficiency in T-cell function, and use of VIG must be under IND application [59].

9. Conclusions

The public health importance of monkeypox disease cannot be underestimated, as the potential for person-to-person transmission is an issue not only among household residents but also among caregivers of sick individuals. Also, the monkeypox virus has the potential to spread via zoonotic reservoirs.

10. References

- [1] Centers for Disease Control and Prevention. Monkeypox-2024 Available from: <https://www.cdc.gov/poxvirus/monkeypox/index.html>
- [2] Durski KN, McCollum AM, Nakazawa Y, et al. Emergence of monkeypox in West Africa and Central Africa, 1970–2017. *Wkly Epidemiol Rec.* 2018;93(11):125–132.
- [3] Monkeypox virus infection in the united states and other non- endemic countries—2022. <https://emergency.cdc.gov/han/2022/han00466.asp>.
- [4] Guarner J, Del Rio C, Malani PN. Monkeypox in 2022—what clinicians need to know [published online ahead of print. *JAMA.* 2022. <https://doi.org/10.1001/jama.2022.10802>
- [5] Von Magnus P, Andersen EK, Petersen KB, Birch Andersen A. A pox-like disease in cynomolgus monkeys. *Acta Pathol Microbiol Scand* 1959; 46: 156–76.
- [6] Breman JG, Kalisa R, Steniowski MV, Zanotto E, Gromyko AI, Arita I. Human monkeypox, 1970–79. *Bull World Health Organ.* 1980; 58(2):165–182.

-
- [7] Monkeypox Outbreak Global Map | monkeypox | Poxvirus | CDC 2023. <https://www.cdc.gov/poxvirus/mpox/> 2023 June.
- [8] Sabeena S. The changing epidemiology of monkeypox and preventive measures: an update. *Arch Virol* 2023; 168:31.
- [9] Jezek Z, Grab B, Szczeniowski M, Paluku KM, Mutombo M. Clinico-epidemiological features of mon- keypox patients with an animal or human source of infection. *Bull World Health Organ.* 1988; 66 (4):459–464.
- [10] Vora S, Damon I and Fulginiti V. Severe eczema vaccinatum in a household contact of a smallpox vaccinee. *Clin Infect Dis.* 2008.
- [11] Burnett JC, Henchal EA, Schmaljohn AL, Bavari S. The evolving field of biodefence: therapeutic developments and diagnostics. *Nat Rev Drug Discov.* 2005; 4:281–97.
- [12] Nolen LD, Osadebe L, Katomba J, Likofata J, Mukadi D, Monroe B, et al. Extended human-to-human transmission during a monkeypox outbreak in the Democratic Republic of the Congo. *Emerg Infect Dis.* 2016; 22(6):1014–1021.
- [13] Olson VA, Laue T, Laker MT, et al. Real-time PCR system for detection of orthopoxviruses and simultaneous identification of smallpox virus. *J Clin Microbiol* 2004; 42:1940–6.
- [14] Chittick G, Morrison M, Brundage T, Nichols WG. Short-term clinical safety profile of brincidofovir: a favorable benefit–risk proposition in the treatment of smallpox. *Antiviral Res.* 2017. <https://doi.org/10.1016/j.antiviral.2017.01.009>
- [15] Bunge EM, Hoet B, Chen L, et al. The changing epidemiology of human mon- keypox-a potential threat? A systematic review. *PLoS Negl Trop Dis* 2022;16(2): e0010141.
- [16] Beer EM, Rao VB. A systematic review of the epidemiology of human monkey- pox outbreaks and implications for out- break strategy. *PLoS Negl Trop Dis* 2019; 13(10):e0007791.
- [17] Yinka-Ogunleye A, Aruna O, Dalhat M, et al. Outbreak of human monkeypox in Nigeria in 2017-18: a clinical and epidemiological report. *Lancet Infect Dis* 2019; 19:872-9
- [18] Doshi RH, Alfonso VH, Morier D, et al. Monkeypox rash severity and animal ex- posures in the Democratic Republic of the Congo. *Ecohealth* 2020;17:64-73.
- [19] Besombes C, Gonofio E, Konamna X, et al. Intrafamily transmission of mon- keypox virus, Central African Republic, 2018. *Emerg Infect Dis* 2019;25:1602-4.
- [20] Vandebogaert M, KwasiBorski A, Gonofio E, et al. Nanopore sequencing of a monkeypox virus strain isolated from a pustular lesion in the Central African Re- public. *Sci Rep* 2022;12:10768.
- [21] Nolen LD, Osadebe L and Katomba J. Extended human-to-human transmission during a monkeypox outbreak in the Democratic Republic of the Congo. *Emerg Infect Dis* 2016;22:1014-21.
- [22] Ogoina, D.; Iroezindu, M.; James, H.I.; Oladokun, R.; Yinka-Ogunleye, A.; Wakama, P.; Otiike-Odibi, B.; Usman, L.M.; Obazee, E and Aruna, O. Clinical Course and Outcome of Human Monkeypox in Nigeria. *Clin. Infect. Dis.* 2020, 71, e210–e214.

- [23] Jezek, Z.; Szczeniowski, M.; Paluku, K.M.; Mutombo, M. Human Monkeypox: Clinical Features of 282 Patients. *J. Infect. Dis.* 1987;156: 293–298.
- [24] WHO. Monkeypox. Available online: <https://www.who.int/news-room/factsheets/detail/monkeypox>. 2022 July.
- [25] Adler, H.; Gould, S.; Hine, P.; Snell, L.B.; Wong, W.; Houlihan, C.F.; Osborne, J.C.; Rampling, T.; Beadsworth, M.B and Duncan, C.J. Clinical features and management of human monkeypox: A retrospective observational study in the UK. *Lancet Infect. Dis.* 2022.
- [26] Osadebe, L.; Hughes, C.M.; Lushima, R.S.; Kabamba, J.; Nguete, B.; Malekani, J.; Pukuta, E.; Karhemere, S.; Tamfum, J.-J.M.; Okitolonda, E.W.; et al. Enhancing case definitions for surveillance of human monkeypox in the Democratic Republic of Congo. *PLOS Negl. Trop. Dis.* 2017, 11, e0005857
- [27] Huhn, G.D.; Bauer, A.M.; Yorita, K.; Graham, M.B.; Sejvar, J.; Likos, A.; Damon, I.K.; Reynolds, M.; Kuehnert, M.J. Clinical Characteristics of Human Monkeypox, and Risk Factors for Severe Disease. *Clin. Infect. Dis.* 2005; 41: 1742–1751
- [28] Mbala, P.K.; Huggins, J.W.; Riu-Rovira, T.; Ahuka, S.M.; Mulembakani, P.; Rimoin, A.W.; Martin, J.W.; Muyembe, J.-J.T. Maternal and Fetal Outcomes Among Pregnant Women with Human Monkeypox Infection in the Democratic Republic of Congo. *J. Infect. Dis.* 2017;216: 824–828
- [29] Damon, I.K. Status of human monkeypox: Clinical disease, epidemiology and research. *Vaccine* 2011; 29: D54–D59.
- [30] Hussein, F. K., Mahmoud, A. J., and Yousif, B. J. Estimation of Immunoglobulin A, Immunoglobulin G, and Immunoglobulin M Antibody Levels in Laboratory Mice Balb/c Infected with *Entamoeba histolytica* and Treatment with Aqueous Extracts of *Cyperus rotundus* and *Thymus serpyllum*. *Polytechnic Journal*.2020; 10(1), 21.
- [31] Yi X.M, Lei Y.L, Li M, Li Z, Li S. The monkeypox virus-host interplays. *Cell Insight*. 2024:100185
- [32] Estep RD, Messaoudi I, O'Connor MA, Li H, Sprague J, Barron A, et al. Deletion of the monkeypox virus inhibitor of complement enzymes locus impacts the adaptive immune response to monkeypox virus in a nonhuman primate model of infection. *Journal of virology*. 2011;85(18):9527-42.
- [33] Shchelkunov SN. An increasing danger of zoonotic orthopoxvirus infections. *PLoS pathogens*. 2013;9(12):e1003756.
- [34] Adler H, Gould S, Hine P, Snell LB, Wong W, Houlihan CF, et al. Clinical features and management of human monkeypox: a retrospective observational study in the UK. *The Lancet Infectious Diseases*. 2022;22(8):1153-62
- [35] Kulesh DA, Loveless BM, Norwood D, et al. Monkeypox virus detection in rodents using real-time 3f-minor groove binder TaqMan assays on the Roche Light Cycler. *Lab Invest* 2004; 84:1200–8.
- [36] Bayer-Garner IB. Monkeypox virus: histologic, immunohistochemical and electron-microscopic findings. *J Cutan Pathol* 2005; 32:28–34.

- [37] Gentile M, Gelderblom HR. Rapid viral diagnosis: role of electron microscopy. *New Microbiol* 2005; 28:1–12.
- [38] Damon IK, Esposito JJ. Poxviruses that infect humans. In: Murray PR, Baron EJ, Jorgensen JH, Pfaller MA, Tenover FC, Tenover MC, eds. *Manual of clinical microbiology*. 8th ed. Washington, DC: ASM Press, 2003:1583–91
- [39] Taher, H. M. Levels Estimation of Iron, Zinc and Copper in the Serum of Children Infected with Giardiasis. *Al-Kitab Journal for Pure Sciences*.2024; 8(02), 120-124.
- [40] Lapa S, Mikheev M, Shchelkunov S, et al. Species-level identification of orthopoxviruses with an oligonucleotide microchip. *J Clin Microbiol* 2002; 40:753–7.
- [41] Guarner J, Del Rio C, Malani PN. Monkeypox in 2022-what clinicians need to know. *JAMA*. 2022;328(2):139–40.
- [42] Centers for Disease Control and Prevention. Impact of monkeypox outbreak on select behaviors. 2022. <https://www.cdc.gov/poxvirus/monkeypox/response/2022/amis-select-behaviors.html#:~:text=In%20an%20online%20survey%20of,encounters%2C%20and%2050%25%20reported%20reducing>. Accessed 19 Oct 2022.
- [43] Rimoin AW, Mulembakani PM, Johnston SC, Lloyd Smith JO, Kisalu NK, Kinkela TL, et al. Major increase in human monkeypox incidence 30 years after smallpox vaccination campaigns cease in the Democratic Republic of Congo. *Proc Natl Acad Sci USA*. 2010;107(37):16262–7.
- [44] Centers for Disease Control and Prevention. Monkeypox and smallpox vaccine guidance. 2022. <https://www.cdc.gov/poxvirus/monkeypox/clinicians/smallpox-vaccine.html>. Accessed 26 Sept 2022.
- [45] Amer, F., Khalil, H. E., Elahmady, M., ElBadawy, N. E., Zahran, W. A., Abdelnasser, M. and Tash, R. M. E. Monkeypox: Risks and approaches to prevention. *Journal of infection and public health*.2023; 16(6), 901-910.
- [46] Guarner J, Del Rio C, Malani PN. Monkeypox in 2022-what clinicians need to know. *JAMA*. 2022;328(2):139–40.
- [47] Reynolds MG, McCollum AM, Nguete B, Lushima RS, Petersen BW. Improving the care and treatment of monkeypox patients in low-resource settings: applying evidence from contemporary bio- medical and smallpox biodefense research. *Viruses*. 2017. <https://doi.org/10.3390/v9120380>.
- [48] Adler H, Gould S, Hine P, et al. Clinical features and management of human monkeypox: a retrospective observational study in the UK. *Lancet Infect Dis*. 2022;S1473–3099(22)00228–6. [https://doi.org/10.1016/S1473-3099\(22\)00228-6](https://doi.org/10.1016/S1473-3099(22)00228-6).
- [49] TPOXX (tecovirimat) [Package Insert]. Corvallis, OR: SIGA Technologies, Inc. 2018. https://www.accessdata.fda.gov/drugs_atfda_docs/label/2018/208627s000lbl.pdf. Accessed 2022 May.
- [50] Russo AT, Grosenbach DW and Chinsangaram J. An overview of tecovirimat for smallpox treatment and expanded anti-ortho- poxvirus applications. *Expert Rev Anti Infect Ther*. 2021. <https://doi.org/10.1080/14787210.2020.1819791>.

- [51] Grosenbach DW, Honeychurch K, Rose EA, et al. Oral tecovirimat for the treatment of smallpox. *N Engl J Med*. 2018. [https://doi.org/ 10.1056/nejmoa1705688](https://doi.org/10.1056/nejmoa1705688).
- [52] Quenelle DC, Buller RML, Parker S, et al. Efficacy of delayed treatment with ST-246 given orally against systemic orthopox- virus infections in mice. *Antimicrob Agents Chemother*. 2007. <https://doi.org/10.1128/AAC.00879-06>.
- [53] Marcinak J, Vora S, Weber S, et al. Household transmission of vaccinia virus from contact with a military smallpox vaccine Illinois and Indiana. *Morb Mortal Wkly Rep*. 2007.
- [54] Taher, H. M. The effect of aqueous and alcoholic extract of *Opuntia ficus indica* on biochemical parameters and kidney function in rats infected with Echinococcosis. *GSC Biological and Pharmaceutical Sciences*.2023; 25(2), 244-248.
- [55] Hahne S, Macey J, Binnendijk RV, et al. Progressive vaccinia in a military smallpox vaccine—United States. *Pediatr Infect Dis J*. 2009. <https://doi.org/10.1097/inf.0b013e3181b18ed0>
- [56] US Food and Drug Administration: FDA approves drug to treat smallpox. <https://www.fda.gov/drugs/drug-safety-and-availability/fda-approves-drug-treat-smallpox>. Accessed 2022 May.
- [57] Bakr, M.M., Taher, H. M. & Mohamed, A. H. The Effect of *Entamoeba Histolytica* Infection on Levels of Adiponectin and Histamine in Children. *Bahrain Medical Bulletin*.2022; 44(2).
- [58] Lanier R, Trost L, Tippin T, et al. Development of CMX001 for the treatment of poxvirus infections. *Viruses*. 2010. <https://doi.org/10.3390/v2122740>.
- [59] Rice AD, Adams MM, Wallace G, et al. Efficacy of CMX001 as a post exposure antiviral in New Zealand white rabbits infected with rabbit pox virus, a model for orthopoxvirus infections of humans. *Viruses*. 2011. <https://doi.org/10.3390/v3010047>.