Review

Male genital damage in COVID-19 patients: Are available data relevant?

Youssef Kharbach*, Abdelhak Khallouk

Urology Department, Tangier University Hospital, Faculty of Medicine, Abdelmalek Essaâdi University, 90000, Tangier, Morocco

Received 25 April 2020; received in revised form 21 May 2020; accepted 10 June 2020
Available online 21 June 2020

KEYWORDS
Coronavirus disease 2019; Severe acute respiratory syndrome coronavirus 2; Male genital system; Testis; Male infertility

Abstract Over the past few weeks, we have observed increasing concern about the possible impact of coronavirus disease 2019 (COVID-19) which is caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2 virus) on male fertility. Precise mechanisms of male reproductive damages are still unclear, but it seems that high temperature resulting from persistent fever and triggering a secondary autoimmune response leading to an autoimmune orchitis are the most likely involved mechanisms. Also, angiotensin conversion enzyme 2 (ACE2) plays a highly important role in cellular entry for SARS-CoV-2 and male genital system presents high ACE2 expression. All these preliminary findings suggest that COVID-19 could impact men's reproductive health. Thus, we examined available data including published and unpublished articles to assess the potential risk of COVID-19 in particular on the male reproductive system.

© 2021 Editorial Office of Asian Journal of Urology. Production and hosting by Elsevier B.V. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

1. Introduction

Many viruses such as mumps virus, hepatitis viruses B/C, herpes simplex virus, human papillomavirus, coxsackie virus, influenza, human immunodeficiency virus (HIV) and severe acute respiratory syndrome coronavirus 1 (SARS-CoV-1) could infect the male genital system and impair fertility [1,2]. Over the past few weeks, we have observed increasing concern about the impact of coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) on male fertility. Thus, we examined available data including published and unpublished articles to assess the potential risk of COVID-19 in particular on the male reproductive system.

2. Viral threat to male fertility

It is known that viruses could infect the testis directly [2]. Also, the male reproductive system has an immunosuppressive environment due to the blood–testis barrier which...
might protect seminal viruses from immune surveillance [1]. For example, mumps viruses has high tropism for the testes; orchitis develops in 20%–30% of cases and could destroy testicular parenchyma [1,2]. Main mechanisms of testis damage in mumps infection are unknown and remain to be clarified [1,3]. However, the most common hypothesis is that testis degeneration is attributed to an increase in testicular temperature as an indirect effect of the inflammatory milieu [1].

Also, Xu et al. [2] conducted a study on autopsy specimens of testis that were obtained from six patients who died of SARS-CoV-1 infection (SARS-CoV-2 is closely related to SARS-CoV-1 with above 85% identity [4]). All six cases had orchitis. Xu et al. [2] noticed a leukocyte infiltration in the SARS-CoV-1 testis that could affect the function of Leydig cells, damage the blood-testis barrier, and destroy directly the seminiferous epithelium. Furthermore, symptoms of orchitis in these patients were not observed or reported clinically.

Precise mechanisms of male reproductive damages in viral infection are still unclear [1], but it seems that high temperature resulting from persistent fever and triggering a secondary autoimmune response leading to an autoimmune orchitis are the most likely mechanisms of male reproductive system damage [1–3].

All this data suggest that male urogenital system may be a potential target of SARS-CoV-2.

3. Epidemiology

SARS-CoV-2 is the virus causing the disease named COVID-19 [5]. This virus is the seventh infecting humans member of the coronorvinae subfamily which are enveloped, single positive-strand RNA viruses [5]. It is listed in the World Health Organization (WHO) Blueprint list for priority pathogens for research due to its epidemic potential [5].

It is also the most contagious [6], and is now declared as a Public Health Emergency of International Concern by the WHO [7]. As of 11 Jun 2020, there have been 7 221 717 confirmed cases with 411 818 deaths [7]. Interestingly, COVID-19 has a lethality that continuously rises with age unlike other respiratory diseases that have a U-shaped lethality curve [5].

Limited data are available at this stage of the pandemic, concerning the involvement of male genital system in COVID-19 patients.

4. Clinical characteristics

Almost 58.1% of patients are males with a median age of 47–49 years and increased severity in the elderly [6,8,9]. Also, the complete clinical manifestation is not clear yet [8]. COVID-19 patients present a wide spectrum of clinical features [5]. They commonly report fever, respiratory symptoms, myalgia or fatigue [8,9]. Most patients present cytokine storm that might cause immune damage [10]. This uncontrolled reaction of the immune system can damage most vital organs and lead to acute kidney injury, various degrees of liver damage, acute cardiac injury, and multiple organ failure [10]. Previous exposure to non-SARS-CoVs in China could explain the disparity between the severity of cases observed in China and those outside China [5]. Male genital system involvement is not clear yet.

5. Pathogenesis: How could SARS-CoV-2 infect male genital system?

Angiotensin conversion enzyme (ACE) converts angiotensin I to angiotensin II in the renin-angiotensin-aldosterone system (RAAS) [11]. ACE2 is an endogenous counter-regulator of classic ACE system that deactivates the angiotensin II [11].

First, the SARS-CoV-2 enters into human cells using ACE2 as receptor which have very similar spike protein three-dimensional (3D) structures with strong binding affinity [12]. ACE2 is present with relatively high expression in respiratory epithelial cells, alveolar cells type I and II, oral cavity, kidney, testis, and intestines [11]. Thus, human cells with ACE2 expression are probably targets of SARS-CoV-2 infection [13]. However, it should still be noted that SARS-CoV-2 invasion is not just about ACE2 and there is probably other vulnerable organs [13]. It seems that high temperature resulting from persistent fever and triggering a secondary autoimmune response leading to an autoimmune orchitis are the most likely involved mechanisms.

6. SARS-CoV-2 and male genital system

Preliminary findings from studies that are, unfortunately, not peer-reviewed and published as preprints, are linking SARS-CoV-2 to male infertility [14–16]. But, it seems to us that it is premature to make definitive conclusions at present.

Ma et al. [14] conducted a study on 81 reproductive-aged men with SARS-CoV-2 infection and 100 age-matched healthy men. They found significantly elevated luteinizing hormone (LH) level, but decreased testosterone/LH and follicle-stimulating hormone (FSH)/LH ratios levels in 81 men with COVID-19 which is suggesting potential hypogonadism. But it seems to us that these findings cannot affirm whether this alteration is due to a direct or indirect effect of SARS-CoV-2 on the testes. Previous studies reported that high temperature contributed to germ cell destruction [2], and high fever in COVID-19 might have an indirect effect on testicular dysfunction. Furthermore, LH release is pulsatile and shows circadian rhythm. We think that there is a biased sampling in Ma et al. study [14]. Actually, COVID-19 patients and control group should be tested at the same time of the day, otherwise, widely varying average levels of LH will be observed.

Also, Fan et al. [15] and Shen et al. [16] found high ACE2 expression in testis (both germ cells and somatic cells) suggesting potential tropism of SARS-CoV-2 to testicular tissues. Shen et al. [16] performed bioinformatics tools to analyze scRNA-seq data of testis for three normal young males, two adult normal males, seven obstructive azoospermia and one nonobstructive azoospermia. They showed that the expression level of ACE2 was related to the age and the peak of positive rate was at 30 years old [16]. These findings are confirmed by the interesting study of Liu et al.
Critical revision of the manuscript warranted. Investigations of the potential male genital damage are evident in patients, as most of them are males [8,9]. Further systematic efforts are needed to elucidate the possible impact of COVID-19 on the male reproductive system. This is particularly important for COVID-19 patients with male genital involvement. In conclusion, the available data are relevant and show that COVID-19 could impact men’s reproductive health inducing spermatogenic failure. In conclusion, the available data are relevant and show that COVID-19 could impact men’s reproductive health inducing spermatogenic failure.

Author contributions

Study concept and design: Youssef Kharbach, Abdelhak Khallouk.

Data acquisition: Youssef Kharbach.

Data analysis: Youssef Kharbach, Abdelhak Khallouk.

Drafting of manuscript: Youssef Kharbach.

Critical revision of the manuscript: Abdelhak Khallouk.

Conflicts of interest

The authors declare no conflict of interest.

References

[1] Liu W, Han R, Wu H, Han D. Viral threat to male fertility. Andrology 2018;50:e13140. https://doi.org/10.1111/and.13140.

[2] Xu J, Qi L, Chi X, Yang J, Wei X, Gong E, et al. Orchitis: a complication of severe acute respiratory syndrome (SARS). Biol Reprod 2006;74:410–6.

[3] Dejucq N, Jéguo B. Viruses in the mammalian male genital tract and their effects on the reproductive system. Microbiol Mol Biol Rev 2001;65:208–31.

[4] Gralinski LE, Menachery VD. Return of the coronavirus: 2019-nCoV. Viruses 2020;12:135. https://doi.org/10.3390/v12020135.

[5] Raoult D, Zumla A, Locatelli F, Ippolito G, Kroemer G. Coronavirus infections: epidemiological, clinical and immunological features and hypotheses. Cell Stress 2020;4:66–75.

[6] Wang C, Horby PW, Hayden FG, Gao GF. A novel coronavirus outbreak of global health concern. Lancet 2020;395:470–3.

[7] WHO. Coronavirus disease (COVID-19) pandemic. https://www.who.int/emergencies/diseases/novel-coronavirus-2019. [Accessed 11 June 2020].

[8] Adhikari SP, Meng S, Wu Y, Mao Y, Ye R, Wang Q, et al. Epidemiology, causes, clinical manifestation and diagnosis, prevention and control of coronavirus disease (COVID-19) during the early outbreak period: a scoping review. Infect Dis Poverty 2020;9:29. https://doi.org/10.1186/s40249-020-00646-x.

[9] Guan W, Ni Z, Hu Y, Liang W, Ou C, He J, et al. Clinical characteristics of coronavirus disease 2019 in China. N Engl J Med 2020;382:1708–20.

[10] Wang Y, Wang Y, Chen Y, Qin Q. Unique epidemiological and clinical features of the emerging 2019 novel coronavirus pneumonia (COVID-19) implicate special control measures. J Med Virol 2020;92:568–76. https://doi.org/10.1002/jmv.25748.

[11] Tolouian R, Vahed SZ, Ghiyasvand S, Tolouian A, Ardalan MA. COVID-19 interactions with angiotensin-converting enzyme 2 (ACE2) and the kinin system; looking at a potential treatment. J Ren Inj Prev 2020;9:e19. https://doi.org/10.34172/jrip.2020.19.

[12] Xu X, Chen P, Wang J, Feng J, Zhou H, Li X, et al. Evolution of the novel coronavirus from the ongoing Wuhan outbreak and modeling of its spike protein for risk of human transmission. Sci China Life Sci 2020;63:457–60.

[13] Zou X, Chen K, Zou J, Han P, Hao J, Han Z. Single-cell RNA-seq data analysis on the receptor ACE2 expression reveals the potential risk of different human organs vulnerable to 2019-nCoV infection. Front Med 2020;14:185–92.

[14] Ma L, Xie W, Li D, Shi L, Mao Y, Xiong Y, et al. Effect of SARS-CoV-2 infection upon male gonadal function: a single center-based study. MedRxiv 2020. https://doi.org/10.1101/2020.03.21.20037267.

[15] Fan C, Li K, Ding Y, Lu W, Wang J. ACE2 expression in kidney and testis may cause kidney and testis damage after 2019-nCoV infection. MedRxiv 2020. https://doi.org/10.1101/2020.02.12.20022418.

[16] Shen Q, Xiao X, Aierken A, Yue W, Wu X, Liao M, et al. The ACE2 expression in Sertoli cells and germ cells may cause male reproductive disorder after SARS-CoV-2 infection. J Cell Mol Med 2020;24:9472–7.

[17] Liu X, Chen Y, Tang W, Zhang L, Chen W, Yan Z, et al. Single-cell transcriptome analysis of the novel coronavirus (SARS-CoV-2) associated gene ACE2 expression in normal and non-obstructive azoospermia (NOA) human male testes. Sci China Life Sci 2020;30:1–10.

[18] Ding Y, He L, Zhang Q, Huang Z, Che X, Hou J, et al. Organ distribution of severe acute respiratory syndrome (SARS) associated coronavirus (SARS-CoV) in SARS patients: implications for pathogenesis and virus transmission pathways. J Pathol 2004;203:622–30.

[19] Ling Y, Xu S, Lin Y, Tian D, Zhu Z, Dai F, et al. Persistence and clearance of viral RNA in 2019 novel coronavirus disease rehabilitation patients. Chin Med J 2020;133:1039–43.