

Risk of COVID-19 in Patients Receiving Anti-VEGF Therapy during the Pandemic

Valtueña Jara^{1*}, López-Galvez MI², Ruiz-Sánchez Daniel¹, Hadi Suhail³ and Duenas Carlos⁴

¹ Department of Dermatology, University Clinical Hospital of Valladolid, Valladolid, Spain

² Department of Ophthalmology, University Clinical Hospital of Valladolid, Valladolid, Spain

³ Department of Dermatology, Icahn School of Medicine at Mount Sinai, New York, NY, Mount Sinai NY Hospital, NY, USA

⁴ Department of Internal Medicine, University Clinical Hospital of Valladolid, Valladolid, Spain

About the Study

Based on observed disease patterns, Corona Virus Disease 2019 (COVID-19) is a systemic multi-organ viral invasion with microvascular injury [1]. Recent studies indicate that COVID-19 could comprise an obliterative micro-angiopathy consisting of endothelitis with altered vessel barrier integrity, promoting angiogenesis and a pro-coagulative state [1,2]. Activation of angiogenic pathways could have an important role in inducing and maintaining the viral vascular effects. Vascular Endothelial Growth Factor (VEGF) is known as the most potent inducing factor to increase angiogenesis and vascular permeability [3]. As a result, vascular endothelial cells could become a potential target to fight against COVID-19. Some studies suggest bevacizumab could be beneficial for patients with severe COVID-19 [4-6]. The role of anti-VEGF agents should be deeper investigated. Risk factors for developing severe COVID-19 is retinal disease and patients being \geq 65 years old [7,8]. Herein, we evaluated the risk of COVID-19 in patients receiving anti-VEGF intravitreal injections during the pandemic period.

The study comprised 431 patients diagnosed with eye conditions (198 men and 233 female), mean age of 76.3 years-old, within one tertiary Spanish Hospital. All patients received intravitreal anti-VEGF therapy (bevacizumab, ranibizumab or aflibercept) in an "as needed" injections regimen. All the patients received, at least, one anti-VEGF intravitreal injection from March to June 2020. Influenza viruses and seasonal coronaviruses and on the viral infection-induced production of inflammatory cytokines.

The patients received an average of 1.47 intravitreal injections from March to June 2020. From the total sample of 431 patients, only 4 patients (3 female and 1 male) tested positive for SARS-CoV-2 (0.86%). Among them, only two had evidence of coronavirus infection and required hospital admission (0.43%), with a mean hospital stage of 8 days. None of them required mechanical ventilation or had other severe complications. No side effects

or other symptoms were reported across all patients and no deaths occurred within that period. Compared to general population data from the same region, the SARS-CoV-2 seroprevalence was much lower in those patients who received anti-VEGF intravitreal injection by comparison with baseline region population (0.86% vs. 7.4% respectively ($P < 0.05$)). Moreover, the prevalence of hospital admission due to COVID-19 was also much lower in those patients treated with intravitreal anti-VEGF agent by comparison with baseline region population (0.43% vs. 4.1% respectively).

Spain is one of the countries severely affected by the ongoing COVID-19 pandemic. The general SARS-CoV-2 seroprevalence of the studied region was 7.4% [9]. On the contrary, in the same region, the studied anti-VEGF treated population had a SARS-CoV-2 seroprevalence of 0.86%. Only 0.43% of the anti-VEGF treated population required hospital admission, despite the fact that these patients had higher SARS-CoV-2 exposure risk due to the need to go to the hospital during the hard-pandemic period. No deaths occurred within the period across anti-VEGF treated patients [10].

Conclusion

Anti-VEGF agents could have an important role in prevention and mitigating the effects of COVID-19. Microvascular endothelial cell injury could activate the clotting pathway. Thus, endothelial cells are increasingly recognized as potential target to prevent thrombotic events and to accelerate thrombus resolution. The possibility that these cells play an important role in COVID-19 should trigger search for future therapeutic opportunities. These observations could help find potential specific pharmacotherapy for fighting against COVID-19 severely ill patients. Further studies are needed.

Conflict of Interest

The authors have no conflicts of interest.

*Address for Correspondence: Dr. Valtueña Jara, Department of Dermatology, Hospital Clínico Universitario de Valladolid, Valladolid, Spain; E-mail: jara.valtueña@gmail.com

Copyright: © 2022 Jara V, et al. This is an open-access article distributed under the terms of the creative commons attribution license which permits unrestricted use, distribution and reproduction in any medium, provided the original author and source are credited.

Received: 30-Jun-2022, Manuscript No. VCRH-22-68173; Editor assigned: 04-Jul-2022, Pre QC No. VCRH-22-68173 (PQ); Reviewed: 18-Jul-2022, QC No. VCRH-22-68173; Revised: 25-Jul-2022, Manuscript No. VCRH-22-68173 (A); Published: 01-Aug-2022, DOI: 10.37421/2736-657X.2022. S2.001.

Funding Sources

None

References

1. Valtuena, Jara, Gerardo Martínez García, Daniel Ruiz Sánchez and María Garayar Cantero, et al. "Vascular Obliteration Because of Endothelial and Myointimal Growth In COVID-19 Patients." *Int J Dermatol* 60(2021): 185-187.
2. Valtuena, Jara, Daniel Ruiz Sánchez, Pilar Manchado López and María Garayar Cantero. "Acral Edema During The COVID-19 Pandemic." *Int J Dermatol* 59(2020): 1155-1157.
3. Apte, Rajendra, Daniel Chen and Napoleone Ferrara. "VEGF in Signaling and Disease: Beyond Discovery and Development." *Cell* 176(2019): 1248-1264.
4. Pang, Jiaojiao, Feng Xu, Gianmarco Aondio and Yu Li, et al. "Efficacy and Tolerability of Bevacizumab in Patients with Severe COVID-19." *Nat Commun* 12(2021): 814.
5. Sahebnasagh, Adeleh, Seyed Mohammad Nabavi, Hamid Reza Khayat Kashani and Safieh Abdollahian, et al. "Anti-VEGF Agents: As Appealing Targets in The Setting Of COVID-19 Treatment In Critically Ill Patients." *Int Immunopharmacol* 101(2021): 108257.
6. Lampropoulou, Dimitra Ioanna, Vanessa Meletia Bala, Eleni Zerva and Evangelia Pliakou, et al. "The Potential Role of The Combined PARP-1 And VEGF Inhibition in Severe SARS-Cov-2 (COVID-19) Infection." *J Chin Med Assoc* 83(2020): 817-821.
7. Korobelnik, Jean François, Anat Loewenstein, Bora Eldem and Antonia Joussem, et al. "Guidance For Anti-VEGF Intravitreal Injections During The COVID-19 Pandemic." *Graefes Arch Clin Exp Ophthalmol* 258(2020): 1149-1156.
8. Boyd, Matt, Daniel Scott, David Squirrel and Graham Wilson. "Proof-of-Concept Calculations to Determine the Health-Adjusted Life-year Trade-off Between Intravitreal anti-VEGF Injections and Transmission of COVID-19." *Clin Exp Ophthalmol* 48(2020): 1276-1285.
9. Ministry of Health, Consumption and Social Welfare of the Government of Spain. National Sero-Epidemiology Study of SARS-CoV-2 Infection in Spain (ENE-Covid). 2020.
10. Prchal, Josef. "Hypoxia And Thrombosis. " *Blood* 132(2018): 348-349.

How to cite this article: Jara,Valtuena,López-Galvez MI , Ruiz-Sánchez Daniel and Hadi Suhail, et al."Risk of COVID-19 in Patients Receiving Anti-VEGF Therapy during the Pandemic". *Virol Curr Res* (6): (S2) (2022) :001