

Partial Human Kluver-Bucy Syndrome Secondary to Right Frontal Arterio-Venous Malformation (AVM) Resection

Fabian Rossi^{1,2*}, Lourdes Benes Lima², Nina Tsakadze^{2,3}, Elisa M Rossi² and Michael Hoffmann^{2,3}

¹Department of Neurology, University of Central Florida College of Medicine, USA

²Department of Neurology, Orlando Veterans Administration Medical Center, USA

³Department of Neurology, University of Central Florida Medical School, USA

ARTICLE INFO

Received Date: January 08, 2020

Accepted Date: January 10, 2020

Published Date: January 13, 2020

KEYWORDS

AVM

Alcohol abuse

Kluver-Bucy

Copyright: © 2020 Fabian Rossi et al., Neurological Disorders & Epilepsy Journal. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Citation for this article: Fabian Rossi, Lourdes Benes Lima, Nina Tsakadze, Elisa M Rossi and Michael Hoffmann. Partial Human Kluver-Bucy Syndrome Secondary to Right Frontal Arterio-Venous Malformation (AVM) Resection. Neurological Disorders & Epilepsy Journal. 2020; 3(1):127

Corresponding author:

Fabian H Rossi,

Department of Neurology, University of Central Florida College of Medicine, Orlando, Florida 32827, USA, Tel: +407-951-1891; Fax:407-513-9317;

Email: fabian.rossi@va.gov

A NEUROIMAGE CASE REPORT

A 67-year-old right-handed male developed apathy, alcohol abuse, episodes of indecent exposure and visual agnosia after right frontal AVM resection. Neuropsychological testing revealed frontal lobe dysfunction. MRI FLAIR imaging revealed signal attenuation in the right frontal lobe (Figures 1A-C). EEG showed no seizure activity. Kluver-Bucy syndrome is a constellation of neuropsychiatric symptoms, classically associated with bilateral temporal damage. Diagnosis of the partial Kluver-Bucy syndrome requires at least 3 of the 6 cardinal features of Kluver-Bucy syndrome, including: placidity, visual agnosia, hypersexuality, hyperorality, hyperphagia, and hypermetamorphosis [1,2]. This case is consistent with the partial Kluver-Bucy syndrome, consequent to unilateral right frontal lesion, a finding not previously reported.

CONFLICT OF INTEREST

Authors report no disclose and no conflict of interests.

REFERENCES

1. Bates GDL and Sturman SG. (1995). Unilateral temporal lobe damage and the partial Kluver-Bucy syndrome. Behavioral Neurology. 8: 103-107.
2. Carroll BT, Goforth HW, Raimonde LA. (2001). Partial Kluver Bucy syndrome: two cases. CNS Spectr. 6: 329-332.

