

# Resolution Of Mysophobia Following Resection Of Large Sphenoid Wing Meningioma

R Elliott, B Rubin, J Jafar

## Citation

R Elliott, B Rubin, J Jafar. *Resolution Of Mysophobia Following Resection Of Large Sphenoid Wing Meningioma*. The Internet Journal of Psychiatry. 2012 Volume 2 Number 1.

## Abstract

Mysophobia is a variant of obsessive-compulsive disorder (OCD) characterized by intense fear of germs, dirt or contamination accompanied by behaviors to decrease contamination and exposure. While brain tumors, traumatic brain injury (TBI) and strokes have been demonstrated in patients with OCD and implicate abnormal function in the left frontotemporal regions, our review found no cases of mysophobia associated with structural lesions. Meningiomas are one of the most common forms of brain tumors and are more frequently diagnosed in women. We describe a woman with a three-year history of severe mysophobia found to have a large, left sphenoid wing meningioma. Following complete surgical resection and improvement in the mass effect on the frontal and temporal lobes, her mysophobia resolved completely and has remained in complete remission for almost 3 years.

## INTRODUCTION

OCD is the fourth most common psychiatric disorder in the United States, affecting 1 to 4% of the population.<sup>9-12</sup> It is characterized by intrusive thoughts (obsessions), which cause anxiety, and repetitive behaviors that reduce that anxiety (compulsions).<sup>12</sup> Mysophobia is an OCD variant characterized by intense fear of germs, dirt or contamination accompanied by behaviors to decrease contamination and exposure such as avoidance of physical contact and frequent hand washing.<sup>1,13</sup> While the pathophysiology of idiopathic OCD and mysophobia is unknown, current theories posit abnormal serotonin activity and altered neuronal activity in the frontal, temporal and striatal regions and their subcortical circuitry.<sup>10-12,14-17</sup> OCD has also been found to occur in patients with brain tumors and following traumatic brain injury (TBI) or stroke—providing converging data suggesting alterations in left frontotemporal brain activity.<sup>2-5,18-22</sup>

Meningiomas are typically benign tumors that arise from the dura mater covering the brain and are one of the most common types of brain tumors. Meningiomas typically present with headache, seizures, focal neurological deficits or personality changes depending on their size and location<sup>6-8,23-26</sup>. To our knowledge, there are no reported cases showing an association between mysophobia and structural lesions of the brain. We describe a woman with a three-year history of severe mysophobia found to have a large, left

sphenoid wing meningioma. Following complete surgical resection and improvement in mass effect on the frontal and temporal lobes, her mysophobia resolved completely.

## CASE REPORT

**History and Examination.** After receiving therapy during a difficult marriage 18 years prior to admission, this 48-year old female was diagnosed as personality disordered, obsessive-compulsive type (OCPD) with generalized anxiety. Benzodiazepines and sertraline provided minimal benefit for her anxiety and obsessive-compulsive symptoms, respectively. Paroxetine, other benzodiazepines and topiramate were tried with varying but overall limited success over the years.

Three years prior to admission, this 48-year-old female developed progressively worsening fears of germs and contamination. She developed significant anxiety from shaking hands with others, eating in restaurants and going to movie theaters. Her symptoms worsened significantly over time and to such an extent that she avoided all physical contact with others, compulsively washed her hands multiple times daily and spending in excess of \$1000 dollars every month on soaps and hand sanitizers. Eventually, her fears caused her to isolate herself and live separately from her family. Other treatments were attempted without success including selegiline, duloxetine, aripiprazole, mirtazipine, n-acetylcysteine (NAC), a glutamate modulator, in an attempt

to reduce her compulsive behaviors. Counter-phobic sensitization offered minimal improvement in her ability to cope with public settings.

Six months prior to admission the patient noted left sided headaches that progressed over time. An MRI scan of the brain with gadolinium contrast demonstrated a large, left sphenoid wing meningioma with significant edema and mass effect on the left frontal and temporal lobes (Figures 1A, B and C). She was neurologically intact on admission but was extremely anxious and visibly uncomfortable with any physical contact during the preoperative neurological examination.

### Figure 1

Figure 1. Gadolinium-enhanced MRI scan of the brain with showing a large, left sphenoid wing meningioma with significant edema and mass effect on the left frontal and temporal lobes (Figure 1A & B). Following complete removal of the meningioma and resolution of mass effect on the brain, this patient's mysophobia resolved (Figure 1C & D).

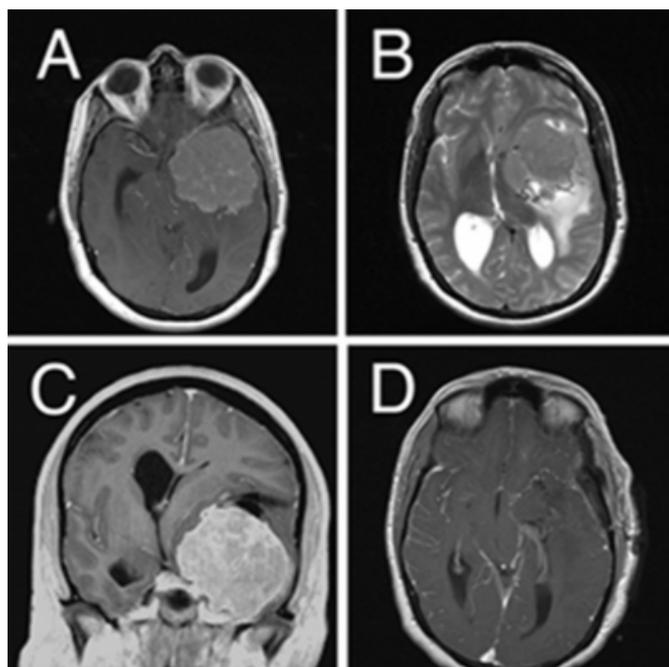


Figure 1. Preoperative MRI shows a large lateral sphenoid wing meningioma causing significant mass effect and edema on the left temporal lobe, insula and frontal operculum (Figure A, B & C). She underwent left pterional craniotomy and gross-total resection of the mass without neurological complication (D).

**Surgical Procedure and Postoperative Course.** She underwent left craniotomy and complete resection of the meningioma with marked resolution of the mass effect and

midline shift (Figure 1D). Postoperatively, she had a third cranial nerve paresis which resolved over a period of 6 weeks. Her mysophobia symptoms resolved rather quickly following surgery. On the first post-operative day, she was able to shake hands with the surgical team with less fear of contamination or the need to immediately wash her hands. At the time of discharge on the fourth postoperative day, she reported no mysophobic symptoms whatsoever. She has remained without symptoms of mysophobia now almost 3 years following surgery and her imaging shows no recurrence of tumor. She has led a normal and active life since discharge following tumor removal and has returned to essentially normal social interaction. She exhibits no anxiety or avoidance behaviors to physical contact or germs nor obsessions concerning contamination.

### DISCUSSION

Originally described by Dr. William Alexander Hammond in 1879,<sup>1</sup> mysophobia is rare variant of obsessive-compulsive disorder (OCD) characterized by intense and pathological fear of dirt, germs or contamination accompanied by behaviors to decrease contamination and exposure.<sup>13</sup> Compulsive hand washing, one of the most common subtypes of OCD, is a prominent feature of this disabling condition. While fear of dirt may underlie the hand washing compulsion of OCD, some contend that it is not primarily a fear of germs per se, as appears to be the case with mysophobia. Rather, patients with hand-washing compulsions in OCD feel that their hands must be washed.<sup>17</sup>

The underlying etiology of OCD and mysophobia is currently unknown. Neurochemically, disruption of serotonin activity is believed to play a prominent role.<sup>10-12, 16</sup> Functional imaging (fMRI, SPECT, PET) studies had identified the frontal, temporal, and striatal regions as possible neuroanatomical substrates of OCD and its variants.<sup>14, 15, 27-33</sup> Using quantitative electroencephalography in patients with OCD, Tot et al. found left frontotemporal dysfunction, especially in women.<sup>34</sup> Others have used magnetoencephalography (MEG) to demonstrate abnormal activity in the left frontotemporal regions,<sup>35</sup> and in the frontotemporal and subcortical/limbic circuitry.<sup>36</sup> Using MRI to measure gray matter volume, other centers have reported a reduced size of the superior temporal gyrus<sup>37</sup> and left orbitofrontal cortex<sup>29, 38</sup> in patients with OCD compared to controls. Numerous centers have also reported on the improvement in the abnormal activity in these regions with successful pharmacological or cognitive-behavioral therapy.<sup>15, 32, 39</sup>

Moreover, patients with brain tumors, strokes and traumatic brain injury (TBI) and OCD have added additional supporting evidence implicating the frontotemporal regions, especially on the left side.<sup>2-5, 19-22, 40, 41</sup> Specifically, Mainio et al. studied levels of obsessive-compulsive symptoms in 59 patients with primary brain tumors and found significantly increased OCD tendencies in patients with left frontal lesions, more so in women.<sup>20</sup>

However, others have suggested that OCD is correlated with greater dysfunction on the right side of brain, but most studies agree that the neuroanatomical substrate may lie in corticostriatal systems, caudate and the orbitofrontal and dorsolateral prefrontal cortices.<sup>12, 15, 33</sup> Whiteside et al. urge caution in the interpretation of functional imaging studies as differences in relative metabolic activity in these different brain regions may not be the underlying cause of the behavioral manifestations.<sup>33</sup>

The primary treatment options for OCD include serotonin reuptake inhibition and psychological treatments such as cognitive behavioral therapy.<sup>12</sup> Selective serotonin reuptake inhibitors (SSRIs) like fluvoxamine and the tricyclic antidepressant clomipramine have been shown to have equal efficacy in patients with OCD and are usually the first-line pharmacological treatment options.<sup>42-45</sup> However, up to 40 to 60% of patients with OCD are refractory to SSRI medications.<sup>46</sup> Limited data suggest that SSRI treatment may be augmented by antipsychotic medications<sup>46, 47</sup> or NAC, an amino acid that modulates glutamate activity.<sup>48, 49</sup>

Brain tumors have been recognized as a cause of psychiatric symptoms since the 16<sup>th</sup> century.<sup>50</sup> Autopsy studies in patients with psychiatric conditions estimate a 1.8 to 3.5% incidence of brain tumors.<sup>24, 50-52</sup> This rate was higher than a control population of patients without psychiatric diagnoses and meningiomas, specifically, constituted a significant proportion of that difference.<sup>51</sup>

Meningiomas are tumors that arise from the meningeal coverings of the brain and are usually benign.<sup>6-8</sup> They account for 13 to 26% of all primary brain tumors and asymptomatic meningiomas are estimated to occur in 1 to 3% of the population.<sup>6, 7</sup> Meningiomas are almost twice as common in females and can present with seizures, focal neurological deficits, or a neuropsychological decline.<sup>8</sup> Moreover, personality, mood and behavioral changes are frequently noted in patients with meningiomas compressing the frontal lobes.<sup>23, 25, 53-56</sup> The primary treatment remains surgical resection,<sup>7, 8</sup> however, minimally invasive options

like focused beam radiosurgery are also used as primary treatment and, specifically, for tumors in areas difficult to treat with microsurgery.<sup>57</sup>

In this report, we described a woman with a large meningioma with significant compression of the left frontotemporal lobes and mysophobia. Given the slow growth rate of these benign tumors, a steady growth over the few years prior to her admission could certainly account for the onset and progression of her mysophobic symptoms. The location of the tumor in the left frontotemporal region is concordant with current data from functional imaging and lesion studies implicating the frontotemporal regions and their subcortical circuitry in the pathophysiology of obsessive thoughts and compulsive behaviors. Removal of the tumor resulted in complete resolution of her mysophobia, providing further evidence that the etiology of the disorder was related to the tumor itself.

Given this patient's long-standing history of anxiety and OCPD and the slow growth rate of meningiomas, early diagnosis in this case was difficult. The refractory nature of her symptoms and the eventual onset of headache prompted imaging. In agreement with others authors,<sup>58</sup> neuroimaging should be considered in patients with new onset psychiatric symptoms, atypical presentations of common psychiatric conditions and cases refractory to standard treatment regimens.

## CONCLUSIONS

This is the first reported case linking a brain tumor to the onset of mysophobia and its resolution following successful surgical treatment. The location of this large meningioma compressing the left frontal and temporal lobes and their subcortical connections converges with neuroanatomical data from patients with idiopathic OCD and OCD in patients with other brain lesions. Neuroimaging should be considered in patients with atypical presentations or refractory cases of psychiatric conditions.

## References

1. Hammond WA. Mysophobia. *Neurological Contributions*; 1879: 40 - 54.
2. Berthier ML, Kulisevsky J, Gironell A, Heras JA. Obsessive-compulsive disorder associated with brain lesions: clinical phenomenology, cognitive function, and anatomic correlates. *Neurology* 1996 Aug;47(2):353-361.
3. Berthier ML, Kulisevsky JJ, Gironell A, Lopez OL. Obsessive-compulsive disorder and traumatic brain injury: behavioral, cognitive, and neuroimaging findings. *Neuropsychiatry Neuropsychol Behav Neurol* 2001 Jan;14(1):23-31.
4. Coetzer BR. Obsessive-compulsive disorder following

- brain injury: a review. *Int J Psychiatry Med* 2004;34(4):363-377.
5. Peterson BS, Bronen RA, Duncan CC. Three cases of symptom change in Tourette's syndrome and obsessive-compulsive disorder associated with paediatric cerebral malignancies. *J Neurol Neurosurg Psychiatry* 1996 Nov;61(5):497-505.
  6. Campbell BA, Jhamb A, Maguire JA, Toyota B, Ma R. Meningiomas in 2009: controversies and future challenges. *Am J Clin Oncol* 2009 Feb;32(1):73-85.
  7. Simpson D. The recurrence of intracranial meningiomas after surgical treatment. *J Neurol Neurosurg Psychiatry* 1957 Feb;20(1):22-39.
  8. Whittle IR, Smith C, Navoo P, Collie D. Meningiomas. *Lancet* 2004 May 8;363(9420):1535-1543.
  9. Rasmussen SA, Eisen JL. The epidemiology and differential diagnosis of obsessive compulsive disorder. *J Clin Psychiatry* 1994 Oct;55 Suppl:5-10; discussion 11-14.
  10. Stein DJ. Advances in the neurobiology of obsessive-compulsive disorder. Implications for conceptualizing putative obsessive-compulsive and spectrum disorders. *Psychiatr Clin North Am* 2000 Sep;23(3):545-562.
  11. Stein DJ. Neurobiology of the obsessive-compulsive spectrum disorders. *Biol Psychiatry* 2000 Feb 15;47(4):296-304.
  12. Stein DJ. Obsessive-compulsive disorder. *Lancet* 2002 Aug 3;360(9330):397-405.
  13. Lawrance G. Mysophobia. *Med J Aust* 1952 Jan 26;1(4):125.
  14. Saxena S, Brody AL, Schwartz JM, Baxter LR. Neuroimaging and frontal-subcortical circuitry in obsessive-compulsive disorder. *Br J Psychiatry Suppl* 1998(35):26-37.
  15. Saxena S, Rauch SL. Functional neuroimaging and the neuroanatomy of obsessive-compulsive disorder. *Psychiatr Clin North Am* 2000 Sep;23(3):563-586.
  16. Stein DJ. Psychobiology of anxiety disorders and obsessive-compulsive spectrum disorders. *CNS Spectr* 2008 Sep;13(9 Suppl 14):23-28.
  17. Sullivan HS. *Clinical Studies in Psychiatry*; 1973.
  18. Griffith JP, Kamthan M. Obsessive-compulsive disorder following closed head injury. *W V Med J* 1998 Jul-Aug;94(4):198-201.
  19. John G, Eapen V, Shaw GK. Frontal glioma presenting as anxiety and obsessions: a case report. *Acta Neurol Scand* 1997 Sep;96(3):194-195.
  20. Mainio A, Hakko H, Niemela A, Salo J, Koivukangas J, Rasanen P. Level of obsessiveness among neurosurgical patients with a primary brain tumor. *J Neuropsychiatry Clin Neurosci* 2005 Summer;17(3):399-404.
  21. Mordecai D, Shaw RJ, Fisher PG, Mittelstadt PA, Guterman T, Donaldson SS. Case study: suprasellar germinoma presenting with psychotic and obsessive-compulsive symptoms. *J Am Acad Child Adolesc Psychiatry* 2000 Jan;39(1):116-119.
  22. Paradis CM, Friedman S, Hatch M, Lazar RM. Obsessive-compulsive disorder onset after removal of a brain tumor. *J Nerv Ment Dis* 1992 Aug;180(8):535-536.
  23. Bassiouni H, Asgari S, Stolke D. Olfactory groove meningiomas: functional outcome in a series treated microsurgically. *Acta Neurochir (Wien)* 2007 Feb;149(2):109-121; discussion 121.
  24. Blackburn I. *Intracranial Tumors Among the Insane*. Washington, DC: Government Printing Office; 1903.
  25. Ciobanu AM, Lisievici MG, Coman TC, et al. Giant wing sphenoid meningioma with principal manifestation depression. *Rom J Morphol Embryol* 2009;50(4):713-717.
  26. Hunter R, Blackwood W, Bull J. Three cases of frontal meningiomas presenting psychiatrically. *Br Med J* 1968 Jul 6;3(5609):9-16.
  27. Adler CM, McDonough-Ryan P, Sax KW, Holland SK, Arndt S, Strakowski SM. fMRI of neuronal activation with symptom provocation in unmedicated patients with obsessive compulsive disorder. *J Psychiatr Res* 2000 Jul-Oct;34(4-5):317-324.
  28. Insel TR. Toward a neuroanatomy of obsessive-compulsive disorder. *Arch Gen Psychiatry* 1992 Sep;49(9):739-744.
  29. Kang DH, Kim JJ, Choi JS, et al. Volumetric investigation of the frontal-subcortical circuitry in patients with obsessive-compulsive disorder. *J Neuropsychiatry Clin Neurosci* 2004 Summer;16(3):342-349.
  30. Kang DH, Kwon JS, Kim JJ, et al. Brain glucose metabolic changes associated with neuropsychological improvements after 4 months of treatment in patients with obsessive-compulsive disorder. *Acta Psychiatr Scand* 2003 Apr;107(4):291-297.
  31. Kwon JS, Kim JJ, Lee DW, et al. Neural correlates of clinical symptoms and cognitive dysfunctions in obsessive-compulsive disorder. *Psychiatry Res* 2003 Jan 20;122(1):37-47.
  32. Nakao T, Nakagawa A, Yoshiura T, et al. Brain activation of patients with obsessive-compulsive disorder during neuropsychological and symptom provocation tasks before and after symptom improvement: a functional magnetic resonance imaging study. *Biol Psychiatry* 2005 Apr 15;57(8):901-910.
  33. Whiteside SP, Port JD, Abramowitz JS. A meta-analysis of functional neuroimaging in obsessive-compulsive disorder. *Psychiatry Res* 2004 Nov 15;132(1):69-79.
  34. Tot S, Ozge A, Comelekoglu U, Yazici K, Bal N. Association of QEEG findings with clinical characteristics of OCD: evidence of left frontotemporal dysfunction. *Can J Psychiatry* 2002 Aug;47(6):538-545.
  35. Maihofner C, Sperling W, Kaltenhauser M, et al. Spontaneous magnetoencephalographic activity in patients with obsessive-compulsive disorder. *Brain Res* 2007 Jan 19;1129(1):200-205.
  36. Amo C, Quesney LF, Ortiz T, et al. Limbic paroxysmal magnetoencephalographic activity in 12 obsessive-compulsive disorder patients: a new diagnostic finding. *J Clin Psychiatry* 2004 Feb;65(2):156-162.
  37. Choi JS, Kim HS, Yoo SY, et al. Morphometric alterations of anterior superior temporal cortex in obsessive-compulsive disorder. *Depress Anxiety* 2006;23(5):290-296.
  38. Choi JS, Kang DH, Kim JJ, et al. Left anterior subregion of orbitofrontal cortex volume reduction and impaired organizational strategies in obsessive-compulsive disorder. *J Psychiatr Res* 2004 Mar-Apr;38(2):193-199.
  39. Schwartz JM, Stoessel PW, Baxter LR, Jr., Martin KM, Phelps ME. Systematic changes in cerebral glucose metabolic rate after successful behavior modification treatment of obsessive-compulsive disorder. *Arch Gen Psychiatry* 1996 Feb;53(2):109-113.
  40. McKeon J, McGuffin P, Robinson P. Obsessive-compulsive neurosis following head injury. A report of four cases. *Br J Psychiatry* 1984 Feb;144:190-192.
  41. Ward CD. Transient feelings of compulsion caused by hemispheric lesions: three cases. *J Neurol Neurosurg Psychiatry* 1988 Feb;51(2):266-268.
  42. Freeman CP, Trimble MR, Deakin JF, Stokes TM, Ashford JJ. Fluvoxamine versus clomipramine in the treatment of obsessive compulsive disorder: a multicenter, randomized, double-blind, parallel group comparison. *J Clin Psychiatry* 1994 Jul;55(7):301-305.
  43. Hollander E. Treatment of obsessive-compulsive spectrum disorders with SSRIs. *Br J Psychiatry Suppl*

1998(35):7-12.

44. Hollander E, Koran LM, Goodman WK, et al. A double-blind, placebo-controlled study of the efficacy and safety of controlled-release fluvoxamine in patients with obsessive-compulsive disorder. *J Clin Psychiatry* 2003 Jun;64(6):640-647.

45. Koran LM, McElroy SL, Davidson JR, Rasmussen SA, Hollander E, Jenike MA. Fluvoxamine versus clomipramine for obsessive-compulsive disorder: a double-blind comparison. *J Clin Psychopharmacol* 1996 Apr;16(2):121-129.

46. Denys D, de Geus F, van Megen HJ, Westenberg HG. A double-blind, randomized, placebo-controlled trial of quetiapine addition in patients with obsessive-compulsive disorder refractory to serotonin reuptake inhibitors. *J Clin Psychiatry* 2004 Aug;65(8):1040-1048.

47. Vulink NC, Denys D, Fluitman SB, Meinardi JC, Westenberg HG. Quetiapine augments the effect of citalopram in non-refractory obsessive-compulsive disorder: a randomized, double-blind, placebo-controlled study of 76 patients. *J Clin Psychiatry* 2009 Jul;70(7):1001-1008.

48. Grant JE, Odlaug BL, Kim SW. N-acetylcysteine, a glutamate modulator, in the treatment of trichotillomania: a double-blind, placebo-controlled study. *Arch Gen Psychiatry* 2009 Jul;66(7):756-763.

49. Lafleur DL, Pittenger C, Kelmendi B, et al. N-acetylcysteine augmentation in serotonin reuptake inhibitor refractory obsessive-compulsive disorder. *Psychopharmacology (Berl)* 2006 Jan;184(2):254-256.

50. Jarquin-Valdivia AA. Psychiatric symptoms and brain

tumors: a brief historical overview. *Arch Neurol* 2004 Nov;61(11):1800-1804.

51. Patton RB, Sheppard JA. Intracranial tumors found at autopsy in mental patients. *Am J Psychiatry* 1956 Oct;113(4):319-324.

52. Raskin N. Intracranial neoplasms in psychotic patients; survey of 2,430 consecutive complete autopsies performed at the Boston State Hospital during the period 1930-1950. *Am J Psychiatry* 1956 Jan;112(7):481-484.

53. Aoki T, Tashiro Y, Fujita K, Kajiwara M, Matsuda Y. [The evaluation of preoperative and postoperative frontal lobe functions in three operative cases of meningioma]. *No Shinkei Geka* 2006 Feb;34(2):161-167.

54. Belyi BI. [Mental disorders in meningiomas of the olfactory fossa]. *Zh Nevropatol Psikhiatr Im S S Korsakova* 1976;76(8):1200-1205.

55. Belyi BI. Mental impairment in unilateral frontal tumours: role of the laterality of the lesion. *Int J Neurosci* 1987 Feb;32(3-4):799-810.

56. Kahn RL, Schlesinger B. Preoperative and postoperative personality changes accompanying frontal lobe meningioma. *J Nerv Ment Dis* 1951 Dec;114(6):492-510.

57. Kondziolka D, Mathieu D, Lunsford LD, et al. Radiosurgery as definitive management of intracranial meningiomas. *Neurosurgery* 2008 Jan;62(1):53-58; discussion 58-60.

58. Madhusoodanan S, Danan D, Moise D. Psychiatric manifestations of brain tumors: diagnostic implications. *Expert Rev Neurother* 2007 Apr;7(4):343-349.

**Author Information**

**Robert E. Elliott, M.D.**  
Neurosurgical Care, LLC

**Benjamin A. Rubin, M.D.**  
Department of Neurosurgery, New York University Langone Medical Center

**Jafar J. Jafar, M.D.**  
Department of Neurosurgery, New York University Langone Medical Center