## It is illegal to post this copyrighted PDF on any website. A Systematic Review of the Utifity of Electroconvulsive Therapy in Broadly Defined Obsessive-Compulsive–Related Disorders

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### ABSTRACT

**Objective:** To assess the efficacy of electroconvulsive therapy (ECT) in *DSM-5* obsessive-compulsive-related disorders (OCRDs) and conditions subsumed under an "extended" OCD spectrum, including tic disorders and self-injurious behaviors.

**Data Sources:** A systematic search of the MEDLINE, Web of Science, Scopus, and LILACS databases and other sources was performed between June 6 and July 2, 2017. Search terms included (Autis\*) AND (ECT OR electroconvulsive), (Self-injur\*) AND (ECT OR electroconvulsive), (Tic\* OR Tourette) AND (ECT OR electroconvulsive), (Body Dysmorphic Disorder OR Dysmorphophobi\*) AND (ECT OR electroconvulsive), (Hoard\*) AND (ECT OR electroconvulsive), (Trichotillomani\*) AND (ECT OR electroconvulsive), (Skin Picking OR Excoriation) AND (ECT OR electroconvulsive), (Grooming) AND (ECT OR electroconvulsive), (Kleptomani\*) AND (ECT OR electroconvulsive), and (Pyromani\*) AND (ECT OR electroconvulsive). No search restrictions (ie, date, language, or document type) were used.

**Study Selection:** Fifty-two records that described the individual responses of OCRDs to ECT (involving 69 patients) were selected.

**Data Extraction:** Clinical data and responses of individual cases were recorded. Data from responders were compared to nonresponders.

**Results:** All records were case reports or case series; there were no randomized controlled trials. Of the 69 OCRD participants who had undergone ECT, a positive response was reported in 73.4% of the cases (including 44.0% of the BDD, 74.1% of the tic disorder, and 85.7% of the self-injurious behavior patients). At follow-up, the majority of responders who had abstained from further ECT had experienced relapse. However, a positive response was obtained in all participants who received a new course of ECT. Patients who responded positively to ECT were likely to report previous unsuccessful treatment with antipsychotics (P<.001) and antidepressants (P=.007).

**Conclusions:** The finding that more than 70% of the reviewed cases showed some response to ECT should not be considered unequivocal evidence of its efficacy in OCRDs. The available evidence suggests that a randomized controlled trial of ECT in OCRDs may be warranted, particularly in severe tic disorders and self-injurious behaviors.

Prim Care Companion CNS Disord 2018;20(5):18r02342

*To cite:* dos Santos-Ribeiro S, de Salles Andrade JB, Quintas JN, et al. A systematic review of the utility of electroconvulsive therapy in broadly defined obsessive-compulsive-related disorders. *Prim Care Companion CNS Disord*. 2018;20(5):18r02342.

To share: https://doi.org/10.4088/PCC.18r02342 © Copyright 2018 Physicians Postgraduate Press, Inc.

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bsessive-compulsive disorder (OCD) is characterized by recurrent and persistent thoughts, urges, or images that are intrusive, unwanted, and distressing (obsessions) and repetitive behaviors or mental acts that the individual performs to ignore, suppress, or neutralize the resulting anxiety or according to rigid rules.<sup>1</sup> OCD affects 2.3% of the general population at some point in their lives and is often chronic and disabling.<sup>2</sup> First-line treatments include high-dose serotonin reuptake inhibitors (SRIs) and exposure and response prevention (ERP).<sup>3</sup> Individuals who show partial response to SRIs are often treated with antipsychotics or, preferentially, outpatient ERP.<sup>4</sup> Up to 10% of patients are generally treatment refractory and are potential candidates for ERP administered in intensive residential treatments<sup>5</sup> or more invasive treatment measures including deep brain stimulation<sup>6</sup> or psychiatric neurosurgery (eg, capsulotomy).<sup>7</sup>

One issue that has often been raised in the literature is whether OCD patients, particularly those who are treatment refractory, will benefit from a course of electroconvulsive therapy (ECT).<sup>8</sup> A systematic review<sup>9</sup> on this topic failed to find evidence supporting the use of ECT in OCD, as the literature was restricted to case reports and series, nonrandomized trials, and cohort studies. Specifically, ECT responders were characterized by a more severe form of OCD with a later onset of OCD symptoms, less depression, and a fewer number of ECT sessions.9 Finally, the finding that ECT responders were less likely to be treated with conventional anti-OCD treatments (ie, adequate trials of SRIs and/or ERP) suggests that ECT could have been avoided if patients were treated more conservatively.9

In addition to insufficient evidence supporting the use of ECT in OCD, it is still unclear whether ECT may be helpful in those affected by obsessivecompulsive-related disorders (OCRDs) other than OCD. In *DSM-5*, OCRDs include body dysmorphic disorder, hoarding disorder, trichotillomania (hair-pulling disorder), and excoriation disorder (skin-picking disorder) among others.<sup>1</sup> Further, despite not being officially recognized as OCRDs in *DSM-5*, some neurodevelopmental disorders have been considered part of an "extended" OCD spectrum

ical Points

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- Despite the lack of randomized controlled trials supporting the efficacy of electroconvulsive therapy in obsessive-compulsive-related disorders, it is still being prescribed to patients with severe forms of repetitive behaviors.
- Maintenance electroconvulsive therapy in obsessivecompulsive-related disorders may be warranted, particularly in cases showing more severe tic disorders and self-injurious behaviors, due to the high relapse rate.

on the basis of clinical and biological commonalities with OCD behaviors, including tic disorders and Tourette's disorder, autism spectrum disorders, and stereotypic movement disorders such as self-injurious behaviors.<sup>10</sup> These neurodevelopmental conditions have been noted to co-occur with each other and to be overrepresented among early onset OCD cases.<sup>11–13</sup>

In contrast to OCD patients, for whom the usefulness of ECT remains elusive, there is good reason to suspect that OCRDs may be responsive to ECT. First, body dysmorphic disorder and hoarding disorder are frequently delusional<sup>14,15</sup> and associated with greater severity of depression.<sup>15,16</sup> Second, symptoms of catatonia (the quintessential "ECTresponsive" neuropsychiatric disorder) overlap substantially with tic disorders and Tourette's disorder<sup>17,18</sup> (eg, echopraxia and echolalia, staring, and grimacing), autism<sup>19-21</sup> (eg, negativism, mutism/verbigeration, and stereotypies/ mannerisms), and self-injurious behaviors<sup>22</sup> (eg, impulsivity and aggressiveness). Also, both cognitive<sup>23,24</sup> and motor OCRDs<sup>25-28</sup> can reach extreme levels of severity, occasionally involving life-threatening behaviors that can result in severe tegumentary lesions,<sup>23</sup> trichobezoars (mass of hairs blocking the bowel),<sup>25</sup> or even sepsis, thus prompting clinicians to consider ECT for some forms of OCRDs.

Despite the reasons for predicting positive responses of OCRD cases to ECT, there also have been negative experiences. For instance, Kalinowsky and Hippius<sup>29</sup> noted that tic disorder patients who "learned" to suppress tics might be unable to do so and deteriorate as a consequence of the detrimental effects of ECT on their ability to focus attention. Worsening of OCD after ECT has been reported in a few cases,<sup>8</sup> but it is unclear whether a similar phenomenon occurs in other OCRDs. To shed light on this poorly understood phenomenon and its treatment, we systematically reviewed studies reporting the effect of ECT in individuals with an official DSM-5 diagnosis of an OCRD, including body dysmorphic disorder, hoarding disorder, trichotillomania, and excoriation disorder, as well as conditions under an "extended" OCD spectrum including tic disorders, autism spectrum disorders, and self-injurious behaviors. This review was based partly on the patient, intervention, comparator, and outcome model<sup>30</sup> and aimed to answer the following questions:

1. Has the effectiveness of ECT in OCRD patients been compared to any sham or active treatment?

sham or active treatment?

- b. If not, what is the highest level of evidence regarding the effectiveness of ECT in OCRD patients?
- 2. How do baseline features of OCRD patients who respond to ECT differ from OCRD patients who do not respond to ECT?

#### **METHODS**

#### **Study Selection Criteria**

*Type of studies.* We searched for randomized controlled trials (RCTs), observational studies, retrospective investigations, case series, and single case reports describing ECT effects on the core repetitive behaviors across the full range of *DSM-5* OCRDs (ie, body dysmorphic disorder, hoarding disorder, trichotillomania, and excoriation disorder), as well as tic disorders, autism spectrum disorders, self-injurious behaviors, kleptomania, and pyromania. Studies that, despite including the targeted populations, reported just the effects of ECT on other syndromes or behaviors (ie, depression in autism spectrum disorders) were preliminarily excluded from our research protocol. Editorials, letters, and reviews were included if sufficiently detailed original data were provided.

*Types of ECT interventions.* We selected all studies reporting the use of modified, unmodified, multiple-monitored, bilateral, and unilateral ECT with any dose, frequency, or level of stimulus. Studies that reported individuals receiving ECT who were also under current or previous pharmacologic treatments (eg, SRIs and antipsychotics) or different forms of psychotherapy (including cognitive-behavioral therapy [CBT] such as ERP, habit reversal, or behavior modification) were also included. Studies describing individuals treated with other brain stimulation techniques such as transcranial magnetic stimulation (TMS), transcranial direct-current stimulation (tDCS), deep brain stimulation (BDS), and vagus nerve stimulation were also included if the patients' response to ECT was also reported.

Type of measures. Given the dearth of instruments employed across and within the target disorders of our review, the primary outcome of interest was the authors' categorical description of clinically meaningful benefits of ECT on repetitive behaviors that characterize DSM-5 OCRDs and the extended OCD spectrum. However, we were also interested in evaluating how frequently reports of ECT use in broadly defined OCRDs described valid instruments for the assessment of severity of repetitive behaviors in each disorder of interest (including, but not restricted to, the Yale-Brown Obsessive-Compulsive Scale [YBOCS] modified for body dysmorphic disorder,<sup>31</sup> the YBOCS modified for neurotic excoriation,<sup>32</sup> the Saving Inventory-Revised,<sup>33</sup> the Massachusetts General Hospital Hairpulling Scale,<sup>34,35</sup> the Yale Global Tic Severity Scale,<sup>36</sup> and the Children's Yale-Brown Obsessive Compulsive Scale modified for autism spectrum disorders<sup>37</sup> among others<sup>38</sup>).

Database	Search Field	Search Terms
Scopus	Article title, abstract, keywords	- (Autis*) AND (ECT OR electroconvulsive)
Web of Science	Торіс	<ul> <li>- (Self-injur*) AND (ECT OR electroconvulsive)</li> </ul>
PubMed	All fields	- (Tic* OR Tourette) AND (ECT OR electroconvulsive)
LILACS	All indexes	- (Body Dysmorphic Disorder OR Dysmorphophobi*) AND (ECT OR electroconvulsive
		<ul> <li>- (Hoard*) AND (ECT OR electroconvulsive)</li> </ul>
		<ul> <li>- (Trichotillomani*) AND (ECT OR electroconvulsive)</li> </ul>
		- (Skin Picking OR Excoriation) AND (ECT OR electroconvulsive)
		- (Grooming) AND (ECT OR electroconvulsive)
		- (Kleptomani*) AND (ECT OR electroconvulsive)
		- (Pyromani*) AND (ECT OR electroconvulsive)

#### Search Methods for Identification of Studies

*Electronic searching.* A systematic search was performed between June 6 and July 2, 2017, in the following databases: MEDLINE, Web of Science, Scopus, and LILACS. The keywords used for the search are listed in Table 1.

**Other sources.** The reference lists of studies that fulfilled inclusion criteria and textbooks deemed relevant for our research question<sup>29,39–45</sup> were also examined for possible new studies. Further, complete and in-progress clinical trials still not published in clinicaltrials.gov and WHO clinical trials registry platforms were also considered for inclusion. No search restrictions (eg, date, language, or document type) were used.

#### **Data Collection and Analysis**

Selection of studies. Repeated titles across databases were removed, and the remaining articles were checked and analyzed individually by our team, which comprised 2 psychiatrists (L.F.F. and J.N.Q.), 2 biomedical scientists (S.S.R. and J.B.S.A.), and 1 psychology student (K.B.B.). During weekly meetings that took place between June 2017 and February 2018, titles and abstracts were analyzed to exclude articles that clearly did not meet the inclusion criteria; then, the full text of selected articles was read for extraction of the data of interest. Eligibility was defined according to inclusion and exclusion criteria for each diagnostic group. When doubts or disagreement between authors regarding the characteristics of the collected variables occurred, a consensus was reached. If the doubts persisted, then corresponding authors of the articles in question were contacted for clarification. If no response was obtained, than the study was excluded from the analysis.

**Data management.** The authors of this review conducted the entire process. If a study fulfilled all of the criteria for inclusion, then its information was entered into a SPSS file (SPSS Inc, Chicago, Illinois). We evaluated experimental and group studies and individual cases. In terms of experimental and group studies, we specifically selected RCTs,



the quality of which we planned to assess according to Cochrane criteria<sup>46</sup> including (1) random sequence generation; (2) allocation concealment; (3) blinding of participants, personnel, and outcome assessors; (4) selective reporting; and (5) incomplete outcome data. In terms of individual cases, whenever possible, demographic and clinical data, alleged indications for ECT use, treatment history before ECT administration, information related to ECT, and clinical efficacy (including follow-up) were collected.

Relevant demographic and clinical data included age, sex, age at OCRD symptom onset, target diagnosis, patterns of comorbidities (eg, OCD/obsessive-compulsive symptoms, other OCRDs, major depressive disorder, bipolar disorder, or schizophrenia), and whether patients were examined with valid assessment tools. Possible indications for ECT were categorized into OCRDs secondary

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Table 2. Description of Demographic, Clinical, and Therapeutic-Related Variables in Main Obsessive-Compulsive–Related Disorder Groups<sup>a</sup>

	Tourette's Disorder	Self-Injurious Behavior	Body Dysmorphic Disorder
Variable	or Tics (n = up to 30)	(n = up to 30)	(n = up to 9)
Age, mean (SD), y	30.8 (15.3)	21.5 (12.6)	31.4 (11.6)
Sex			
Female	11 (36.7)	9 (30)	3 (60)
Male	19 (63.3)	21 (70)	2 (40)
Age at onset of repetitive behavior, mean (SD), y	12.6 (14.6)	12.3 (9.3)	26 (19)
Course of main repetitive behavior disorder			
Chronic	24 (96)	20 (87)	5 (100)
Episodic	1 (4)	3 (13)	0
Major comorbidities			
Obsessive-compulsive symptoms	20 (66.7)	5 (16.7)	1 (20)
Major depressive disorder	12 (40)	8 (27.6)	5 (100)
Autism	8 (26.7)	15 (50)	0
Bipolar disorder	3 (10)	8 (26.7)	0
Schizophrenia-related disorders	2 (6.7)	8 (26.7)	0
Treatment history			
Antipsychotic prescribed	24 (80)	23 (76.7)	3 (33.3)
SRI prescribed	17 (56.7)	17 (56.7)	3 (33.3)
Psychotherapy	7 (23.3)	18 (60)	1 (11.1)
CBT	3 (10)	15 (50)	1 (11.1)
No. of ECT indications			
>1	20 (66.7)	25 (83.3)	1 (11.1)
1	5 (16.7)	2 (6.7)	2 (22.2)

<sup>a</sup>Values shown as n (%) unless otherwise noted.

Abbreviations: CBT = cognitive-behavioral therapy, ECT = electroconvulsive therapy, SRI = serotonin reuptake inhibitor.

to other conditions known to be treatable with ECT (eg, depression); treatment-resistant OCRDs; severe OCRDs (ie, conditions resulting in self-injury or risks for patients' physical health); severe major depressive disorder with suicidality or psychosis; catatonia; severe mania, psychosis, or agitation; previous response to ECT or seizures; and the presence of drug side effects or other indications.

We also collected data on individuals' treatment history including previous use of SRIs, antipsychotics, any form of psychotherapy, and concomitant use of antidepressants and psychotherapy during ECT. Information on ECT parameters such as type of electrode placement (unilateral or bilateral), total number of ECT sessions, and weekly frequency of administration was also registered. Finally, data related to patterns of ECT response were collected, including positive versus negative response of OCRDs or associated depression to ECT, whether this response was based on a valid assessment tool or simply the authors' subjective clinical impression, whether patients have been followed up (reassessed) in the absence of ECT use, whether there was a relapse during follow-up, whether a second course of ECT was required, and, if a second course of ECT was required, whether the patient responded.

Statistical analysis. As previously reported,<sup>9</sup> a priori analyses of ECT responders versus nonresponders were planned on the basis of the following variables: age, sex, age at OCRD symptom onset, main OCRD, course of illness, major comorbidities, previous exposure to SRIs or antipsychotics, previous psychotherapy (including CBT), indication for ECT, and number of different indications for ECT. Categorical variables were described in frequencies and compared using  $\chi^2$  square and Fisher exact test, and continuous variables were compared using the Mann-Whitney *U* test. Correction for multiple comparisons was performed using the Benjamini-Hochberg test<sup>47</sup> with a false discovery rate set at 0.25.

#### RESULTS

The search resulted in 833 articles, of which 384 represented repeated references. After initial selection, the titles and abstracts of the remaining 449 articles were further screened, leaving 95 articles for eligibility assessment according to the selection criteria. Eventually, 57<sup>18,45,48-102</sup> articles included OCRD individuals treated with ECT allowing qualitative analysis, and  $52^{18,45,48-97}$ articles included information allowing quantitative analysis. The studies by Ghaziuddin et al,<sup>71</sup> Yero et al,<sup>98</sup> Larkin,<sup>99</sup> Bailine and Petraviciute,<sup>100</sup> and Thuppal and Fink<sup>101</sup> were excluded from the quantitative analysis for lacking report of the effects of ECT on repetitive behaviors. We also included studies that did not focus primarily on ECT response but that, nevertheless, allowed collection of data related to response of OCRDs to ECT.<sup>48-53</sup> Figure 1 shows the PRISMA diagram of the research steps.

We were unable to find any RCTs, controlled before and after studies, interrupted time series studies, historically controlled studies, cohort studies, case control studies, or cross-sectional studies. All identified articles were case reports or series. Supplementary Table 1 lists the included studies, their design and number of enrolled participants, and the primary indication for ECT administration. A total of 69 OCRD individuals were treated with ECT, of whom 69.2% presented self-injurious behaviors of different etiologies (autism spectrum disorders, tic disorders or Tourette's disorder, and delusions), 35.4% Tourette's disorder

#### It is illegal to post this copyrighted PDF on any website. Table 3. Comparison Between Sociodemographic, Clinical, and Therapeutic Features of Patients Who Responded Versus Those Who Did Not Respond to ECT<sup>a</sup>

VariableResponded (n=up to 47)Did Not Respond (n=up to 17)StatisticsVariable $(n=up to 47)$ $(n=up to 17)$ StatisticsAge, mean (SD), y $25.5$ (13.2) $27.2$ (14.8) $Z=-0.431, P=.66$ Sex $P=.13^*$ Female15 (31.9)7 (53.8)Male $32$ (68.1)6 (46.2)Age at onset of repetitive behavior, mean (SD), y14.4 (14.4)13 (10.6) $Z=-0.098, P=.92$ Main repetitive behavior disorder $23$ (48.9) $5$ (29.4) $P=.08^*$ $P=.08^*$ Self-injurious behavior23 (48.9) $5$ (29.4) $P=.21^*$ $P=.21^*$ Tourette's disorder or tics $20$ (42.6) $7$ (41.2) $P=.21^*$ Body dysmorphic disorder $4$ (8.5) $5$ (29.4) $P=.21^*$ Chronic $36$ (94.7) $9$ (81.8) $P=.21^*$ Episodic $2$ (5.3) $2$ (18.2) $P=.31$ Major depressive disorder $17$ (36.2) $4$ (30.8) $P=.49^*$ Autism $17$ (36.2) $4$ (30.8) $P=.49^*$ Bipolar disorder $8$ (17) $1$ (7.7) $P=.36^*$ Schizophrenia-related disorders $7$ (14.9) $2$ (15.4) $P=.63^*$ Treatment history $1444/7$ $5$ (29.4) $\chi^2_1=1.207, P=.27$ Antipsychotic prescribed40 (85.1) $6$ (35.3) $P<.001^*$ SRI prescribed29 (61.7) $4$ (23.5) $\chi^2_1=7.284, P=.007$ Psychotherapy $21$ (44.7) $5$ (29.4) $\chi^2_1=3.562, P=.05$ Indication for ECTT $7$ (36.2) $2$ (11.8) <t< th=""><th></th><th></th><th></th><th></th></t<>				
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Male       32 (68.1)       6 (46.2)         Age at onset of repetitive behavior, mean (SD), y       14.4 (14.4)       13 (10.6) $Z = -0.098, P = .92$ Main repetitive behavior       23 (48.9)       5 (29.4)         Tourette's disorder or tics       20 (42.6)       7 (41.2)         Body dysmorphic disorder       4 (8.5)       5 (29.4)         Course of main repetitive behavior disorder $P = .21*$ Chronic       36 (94.7)       9 (81.8)         Episodic       2 (5.3)       2 (18.2)         Major comorbidities       0       5 (94.7)         Obsessive-compulsive symptoms       18 (38.3)       7 (53.8) $\chi^2_1 = 1.013, P = .31$ Major depressive disorder       17 (36.2)       5 (41.7) $P = .48*$ Autism       17 (36.2)       4 (30.8) $P = .49*$ Bipolar disorder       8 (17)       1 (7.7) $P = .63*$ Treatment history       7 (14.9)       2 (15.4) $P = .01*$ Antipsychotic prescribed       40 (85.1)       6 (35.3) $P < .001*$ SRI prescribed       29 (61.7)       4 (23.5) $\chi^2_1 = 1.207, P = .27$ CBT       17 (36.2)       2 (11.8) $\chi^3_1 = 3.562, P = .05$	Female	15 (31.9)	7 (53.8)	
Age at onset of repetitive behavior, mean (SD), y       14.4 (14.4)       13 (10.6) $Z = -0.098, P = .92$ Main repetitive behavior disorder       23 (48.9)       5 (29.4)         Self-injurious behavior       23 (48.9)       5 (29.4)         Tourette's disorder or tics       20 (42.6)       7 (41.2)         Body dysmorphic disorder       4 (8.5)       5 (29.4)         Course of main repetitive behavior disorder $P = .21*$ Chronic       36 (94.7)       9 (81.8)         Episodic       2 (5.3)       2 (18.2)         Major comorbidities       0       5 (29.4)         Obsessive-compulsive symptoms       18 (38.3)       7 (53.8) $\chi^2_1 = 1.013, P = .31$ Major depressive disorder       17 (36.2)       5 (41.7) $P = .48^*$ Autism       17 (36.2)       5 (43.0) $P = .49^*$ Bipolar disorder       8 (17)       1 (7.7) $P = .36^*$ Schizophrenia-related disorders       7 (14.9)       2 (15.4) $P = .63^*$ Treatment history       11 (44.7)       5 (29.4) $\chi^2_1 = 1.207, P = .27^*$ Antipsychotic prescribed       40 (85.1)       6 (35.3) $P < .001^*$ SRI prescribed       29 (61.7)       4 (23.5)	Male	32 (68.1)	6 (46.2)	
Main repetitive behavior disorder $P=.08^*$ Self-injurious behavior       23 (48.9)       5 (29.4)         Tourette's disorder or tics       20 (42.6)       7 (41.2)         Body dysmorphic disorder       4 (8.5)       5 (29.4)         Course of main repetitive behavior disorder $P=.21^*$ Chronic       36 (94.7)       9 (81.8)         Episodic       2 (5.3)       2 (18.2)         Major comorbidities       0       5 (29.4)         Obsessive-compulsive symptoms       18 (38.3)       7 (53.8) $\chi^2_1 = 1.013, P=.31$ Major depressive disorder       17 (36.2)       5 (41.7) $P=.48^*$ Autism       17 (36.2)       5 (41.7) $P=.63^*$ Schizophrenia-related disorders       7 (14.9)       2 (15.4) $P=.63^*$ Treatment history       40 (85.1)       6 (35.3) $P <.001^*$ Antipsychotic prescribed       29 (61.7)       4 (23.5) $\chi^2_1 = 7.284, P =.007$ Psychotherapy       21 (44.7)       5 (29.4) $\chi^2_1 = 3.562, P = .05$ Indication for ECT       Treatment resistance       35 (77.8)       6 (85.7) $P =.53^*$ Severe mania, psychosis       35       5 $P =.63^*$ su	Age at onset of repetitive behavior, mean (SD), y	14.4 (14.4)	13 (10.6)	Z=-0.098, P=.92
Self-injurious behavior23 (48.9)5 (29.4)Tourette's disorder or tics20 (42.6)7 (41.2)Body dysmorphic disorder4 (8.5)5 (29.4)Course of main repetitive behavior disorder $P=.21*$ Chronic36 (94.7)9 (81.8)Episodic2 (5.3)2 (18.2)Major comorbidities $2$ (5.3)2 (18.2)Major depressive disorder17 (36.2)5 (41.7)Autism17 (36.2)4 (30.8)P=.49*Bipolar disorder8 (17)I (7.7)P=.36*Schizophrenia-related disorders7 (14.9)2 (15.4)Perescribed40 (85.1)6 (35.3) $P < .001*$ SRI prescribed29 (61.7)4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P = .007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P = .007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECTTTTreatment resistance35 (77.8)6 (85.7)Severe major depressive disorder with14 (31.1)2 (28.6)suicidality or psychosisT2 (28.6) $P = .63*$ Severe major depressive disorder with13 (28.9)2 (28.6)Severe major depressive disorder with14 (31.1) </td <td>Main repetitive behavior disorder</td> <td></td> <td></td> <td>P=.08*</td>	Main repetitive behavior disorder			P=.08*
Tourette's disorder or tics20 (42.6)7 (41.2)Body dysmorphic disorder4 (8.5)5 (29.4)Course of main repetitive behavior disorder $P=.21^*$ Chronic36 (94.7)9 (81.8)Episodic2 (5.3)2 (18.2)Major comorbidities $0$ (5.3) $2$ (18.2)Major depressive disorder17 (36.2)5 (41.7)Autism17 (36.2)4 (30.8)P=.48*8 (17)1 (7.7)P=.36*Schizophrenia-related disorders7 (14.9)2 (15.4)P=.63*Treatment history $21$ (44.7)5 (29.4)Antipsychotic prescribed40 (85.1)6 (35.3)Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P=.27$ CBTCBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P=.05$ Indication for ECTTreatment resistance35 (77.8)6 (85.7)Severe major depressive disorder with14 (31.1)2 (28.6)severe major depressive disorder with14 (31.1)2 (28.6)severe major depressive disorder with13 (28.9)2 (28.6)Severe major depressive disorder with13 (28.9)2 (28.6)severe major symptoms13 (28.9)2 (28.6)Previous response to ECT or seizure1 (2.2)0Previous response to ECT or seizure1 (2.2)0	Self-injurious behavior	23 (48.9)	5 (29.4)	
Body dysmorphic disorder4 (8.5)5 (29.4)Course of main repetitive behavior disorder $P = .21^*$ Chronic36 (94.7)9 (81.8)Episodic2 (5.3)2 (18.2)Major comorbidities $2 (5.3)$ 2 (18.2)Obsessive-compulsive symptoms18 (38.3)7 (53.8) $\chi^2_1 = 1.013, P = .31$ Major depressive disorder17 (36.2)5 (41.7) $P = .48^*$ Autism17 (36.2)4 (30.8) $P = .49^*$ Bipolar disorder8 (17)1 (7.7) $P = .36^*$ Schizophrenia-related disorders7 (14.9)2 (15.4) $P = .63^*$ Treatment history $21 (61.7)$ 4 (23.5) $\chi^2_1 = 7.284, P = .007$ Antipsychotic prescribed40 (85.1)6 (35.3) $P < .001^*$ SRI prescribed29 (61.7)4 (23.5) $\chi^2_1 = -3.562, P = .57$ Indication for ECTTreatment resistance35 (77.8)6 (85.7) $P = .53^*$ Severe repetitive behavior35 (77.8)4 (50) $P = .11^*$ Severe major depressive disorder with14 (31.1)2 (28.6) $P = .68^*$ Drug-related side effects8 (17.8)1 (14.3) $P = .65^*$ Severe maja, psychosis, or agitation15 (33.3)1 (14.3) $P = .65^*$ Severe major depressive disorder5 (11.1)2 (28.6) $P = .38^*$ Drug-related side effects8 (17.8)1 (14.3) $P = .65^*$ Severe major depressive disorder5 (11.1)2 (28.6) $P = .38^*$ Drug-related side effects8 (17.8)1 (14.3)	Tourette's disorder or tics	20 (42.6)	7 (41.2)	
Course of main repetitive behavior disorder $P = 21^*$ Chronic         36 (94.7)         9 (81.8)           Episodic         2 (5.3)         2 (18.2)           Major comorbidities         0         2 (5.3)         2 (18.2)           Major comorbidities         7 (36.2)         5 (41.7) $P = .48^*$ Autism         17 (36.2)         5 (41.7) $P = .48^*$ Autism         17 (36.2)         4 (30.8) $P = .49^*$ Bipolar disorder         8 (17)         1 (7.7) $P = .36^*$ Schizophrenia-related disorders         7 (14.9)         2 (15.4) $P = .63^*$ Treatment history	Body dysmorphic disorder	4 (8.5)	5 (29.4)	
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	Course of main repetitive behavior disorder			P=.21*
Episodic2 (5.3)2 (18.2)Major comorbidities $2(5.3)$ 2 (18.2)Major comorbidities $2(5.3)$ $2(18.2)$ Major comorbidities $2(5.3)$ $2(18.2)$ Major depressive disorder17 (36.2) $5(41.7)$ Autism17 (36.2) $4(30.8)$ $P=.49^*$ Bipolar disorder8 (17) $1(7.7)$ $P=.36^*$ Schizophrenia-related disorders7 (14.9) $2(15.4)$ $P=.63^*$ Treatment history $2(16.7)$ $4(23.5)$ $\chi^2_1 = 7.284, P=.007$ Psychothcic prescribed29 (61.7) $4(23.5)$ $\chi^2_1 = 1.207, P=.27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P=.05$ Indication for ECTTreatment resistance35 (77.8)6 (85.7) $P=.53^*$ Severe repetitive behavior35 (77.8)4 (50) $P=.11^*$ Severe major depressive disorder with14 (31.1)2 (28.6) $P=.63^*$ suicidality or psychosis $3(28.9)$ 2 (28.6) $P=.68^*$ Drug-related side effects8 (17.8)1 (14.3) $P=.29^*$ Catatonia symptoms13 (28.9)2 (28.6) $P=.68^*$ Drug-related side effects8 (17.8)1 (14.3) $P=.65^*$ Secondary repetitive behavior5 (11.1)2 (28.6) $P=.23^*$ No. of ECT indications $P=.06^*$ $P=.06^*$ No. of ECT indications $P=.06^*$ $P=.06^*$	Chronic	36 (94.7)	9 (81.8)	
Major comorbiditiesObsessive-compulsive symptoms18 (38.3)7 (53.8) $\chi^2_1 = 1.013, P = .31$ Major depressive disorder17 (36.2)5 (41.7) $P = .48^*$ Autism17 (36.2)4 (30.8) $P = .49^*$ Bipolar disorder8 (17)1 (7.7) $P = .36^*$ Schizophrenia-related disorders7 (14.9)2 (15.4) $P = .63^*$ Treatment history $V^2_1 = 7.284, P = .007$ $Y^2_1 = 7.284, P = .007$ Antipsychotic prescribed40 (85.1)6 (35.3) $P < .001^*$ SRI prescribed29 (61.7)4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECTTreatment resistance35 (77.8)6 (85.7) $P = .53^*$ Severe repetitive behavior35 (77.8)4 (50) $P = .11^*$ Severe major depressive disorder with14 (31.1)2 (28.6) $P = .63^*$ suicidality or psychosis $S$ $S$ $S$ $S$ Severe mania, psychosis, or agitation15 (33.3)1 (14.3) $P = .29^*$ Catatonia symptoms13 (28.9)2 (28.6) $P = .68^*$ Drug-related side effects8 (17.8)1 (14.3) $P = .65^*$ Secondary repetitive behavior5 (11.1)2 (28.6) $P = .68^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 140 (88.9)4 (57.1)15 (11.1)3 (42.9)	Episodic	2 (5.3)	2 (18.2)	
Obsessive-compulsive symptoms       18 (38.3)       7 (53.8) $\chi^2_1 = 1.013, P = .31$ Major depressive disorder       17 (36.2)       5 (41.7) $P = .48^*$ Autism       17 (36.2)       4 (30.8) $P = .49^*$ Bipolar disorder       8 (17)       1 (7.7) $P = .36^*$ Schizophrenia-related disorders       7 (14.9)       2 (15.4) $P = .63^*$ Treatment history       Treatment history $40 (85.1)$ 6 (35.3) $P < .001^*$ SRI prescribed       29 (61.7)       4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy       21 (44.7)       5 (29.4) $\chi^2_1 = 3.562, P = .05$ Indication for ECT       17 (36.2)       2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECT       5 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) <t< td=""><td>Major comorbidities</td><td></td><td></td><td></td></t<>	Major comorbidities			
Major depressive disorder17 (36.2) $5$ (41.7) $P=.48^*$ Autism17 (36.2)4 (30.8) $P=.49^*$ Bipolar disorder8 (17)1 (7.7) $P=.36^*$ Schizophrenia-related disorders7 (14.9)2 (15.4) $P=.63^*$ Treatment history $10$ (85.1)6 (35.3) $P<.001^*$ Antipsychotic prescribed29 (61.7)4 (23.5) $\chi^2_1 = 7.284, P=.007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P=.27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P=.05$ Indication for ECT $Treatment resistance$ 35 (77.8)6 (85.7)Severe repetitive behavior35 (77.8)6 (85.7) $P=.53^*$ Severe major depressive disorder with14 (31.1)2 (28.6) $P=.63^*$ suicidality or psychosis $Severe$ main, psychosis, or agitation15 (33.3)1 (14.3) $P=.29^*$ Catatonia symptoms13 (28.9)2 (28.6) $P=.68^*$ $P=.06^*$ Drug-related side effects8 (17.8)1 (14.3) $P=.29^*$ Secondary repetitive behavior5 (11.1)2 (28.6) $P=.36^*$ No. of ECT indications $P=.06^*$ $P=.06^*$	Obsessive-compulsive symptoms	18 (38.3)	7 (53.8)	$\chi^2_1 = 1.013, P = .31$
Autism17 (36.2)4 (30.8) $P=.49^*$ Bipolar disorder8 (17)1 (7.7) $P=.36^*$ Schizophrenia-related disorders7 (14.9)2 (15.4) $P=.63^*$ Treatment history740 (85.1)6 (35.3) $P < .001^*$ Antipsychotic prescribed29 (61.7)4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECTTreatment resistance35 (77.8)6 (85.7)Severe repetitive behavior35 (77.8)4 (50) $P = .11^*$ Severe major depressive disorder with14 (31.1)2 (28.6) $P = .63^*$ suicidality or psychosis58 (17.8)1 (14.3) $P = .29^*$ Catatonia symptoms13 (28.9)2 (28.6) $P = .68^*$ Drug-related side effects8 (17.8)1 (14.3) $P = .23^*$ Previous response to ECT or seizure1 (2.2)0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$	Major depressive disorder	17 (36.2)	5 (41.7)	P=.48*
Bipolar disorder8 (17)1 (7.7) $P=.36^*$ Schizophrenia-related disorders7 (14.9)2 (15.4) $P=.63^*$ Treatment historyAntipsychotic prescribed40 (85.1)6 (35.3) $P < .001^*$ SRI prescribed29 (61.7)4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy21 (44.7)5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT17 (36.2)2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECTTreatment resistance35 (77.8)6 (85.7) $P = .53^*$ Severe repetitive behavior35 (77.8)4 (50) $P = .11^*$ Severe major depressive disorder with14 (31.1)2 (28.6) $P = .63^*$ suicidality or psychosisSevere maina, psychosis, or agitation15 (33.3)1 (14.3) $P = .29^*$ Catatonia symptoms13 (28.9)2 (28.6) $P = .68^*$ Drug-related side effects8 (17.8)1 (14.3) $P = .65^*$ Secondary repetitive behavior5 (11.1)2 (28.6) $P = .68^*$ $P = .06^*$ $P = .06^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ $P = .06^*$ $P = .06^*$	Autism	17 (36.2)	4 (30.8)	P=.49*
Schizophrenia-related disorders       7 (14.9)       2 (15.4) $P = .63^*$ Treatment history       Antipsychotic prescribed       40 (85.1)       6 (35.3) $P < .001^*$ SRI prescribed       29 (61.7)       4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy       21 (44.7)       5 (29.4) $\chi^2_1 = 1.207, P = .27$ CBT       17 (36.2)       2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECT       Treatment resistance       35 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^*$ suicidality or psychosis       Severe maia, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$	Bipolar disorder	8 (17)	1 (7.7)	P=.36*
Treatment history       Antipsychotic prescribed       40 (85.1)       6 (35.3) $P < .001^*$ SRI prescribed       29 (61.7)       4 (23.5) $\chi^2_1 = 7.284$ , $P = .007$ Psychotherapy       21 (44.7)       5 (29.4) $\chi^2_1 = 1.207$ , $P = .27$ CBT       17 (36.2)       2 (11.8) $\chi^2_1 = 3.562$ , $P = .05$ Indication for ECT       Treatment resistance       35 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^*$ suicidality or psychosis       Severe mania, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .29^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .68^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1)         1       5 (11.1)       3 (42.9)	Schizophrenia-related disorders	7 (14.9)	2 (15.4)	P=.63*
Antipsychotic prescribed       40 (85.1)       6 (35.3) $P < .001^*$ SRI prescribed       29 (61.7)       4 (23.5) $\chi^2_1 = 7.284, P = .007$ Psychotherapy       21 (44.7)       5 (29.4) $\chi^2_1 = 7.284, P = .007$ CBT       17 (36.2)       2 (11.8) $\chi^2_1 = 3.562, P = .57$ Indication for ECT       Treatment resistance       35 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^*$ suicidality or psychosis       5       77.8)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$	Treatment history			
SRI prescribed       29 (61.7)       4 (23.5) $\chi_{1}^{2} = 7.284, P = .007$ Psychotherapy       21 (44.7)       5 (29.4) $\chi_{1}^{2} = 1.207, P = .27$ CBT       17 (36.2)       2 (11.8) $\chi_{1}^{2} = 3.562, P = .05$ Indication for ECT       7       7.88       6 (85.7) $P = .53^{*}$ Severe repetitive behavior       35 (77.8)       6 (85.7) $P = .53^{*}$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^{*}$ suicidality or psychosis       5       77.8)       1 (14.3) $P = .29^{*}$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^{*}$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^{*}$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .68^{*}$ No. of ECT indications $P = .06^{*}$ $P = .06^{*}$ > 1       40 (88.9)       4 (57.1) $1 (4.2)$	Antipsychotic prescribed	40 (85.1)	6 (35.3)	P<.001*
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	SRI prescribed	29 (61.7)	4 (23.5)	$\chi^2_1 = 7.284, P = .007$
CBT       17 (36.2)       2 (11.8) $\chi^2_1 = 3.562, P = .05$ Indication for ECT       7reatment resistance       35 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^*$ suicidality or psychosis       5       77.8)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1) $1 = .06^*$	Psychotherapy	21 (44.7)	5 (29.4)	$\chi^2_1 = 1.207, P = .27$
Indication for ECT       35 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^*$ suicidality or psychosis       severe mania, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .68^*$ No. of ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1) $1$ 1       5 (11.1)       3 (42.9) $2$	CBT	17 (36.2)	2 (11.8)	$\chi^2_1 = 3.562, P = .05$
Treatment resistance       35 (77.8)       6 (85.7) $P = .53^*$ Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with       14 (31.1)       2 (28.6) $P = .63^*$ suicidality or psychosis       5       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .38^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .38^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1) $14(29)$	Indication for ECT			
Severe repetitive behavior       35 (77.8)       4 (50) $P = .11^*$ Severe major depressive disorder with suicidality or psychosis       14 (31.1)       2 (28.6) $P = .63^*$ Severe mania, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1) $1 (42.9)$	Treatment resistance	35 (77.8)	6 (85.7)	P=.53*
Severe major depressive disorder with suicidality or psychosis       14 (31.1)       2 (28.6) $P = .63^*$ Severe mania, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1)         1       5 (11.1)       3 (42.9)	Severe repetitive behavior	35 (77.8)	4 (50)	P=.11*
suicidality or psychosis         Severe mania, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .65^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1)         1       5 (11.1)       3 (42.9)	Severe major depressive disorder with	14 (31.1)	2 (28.6)	P=.63*
Severe mania, psychosis, or agitation       15 (33.3)       1 (14.3) $P = .29^*$ Catatonia symptoms       13 (28.9)       2 (28.6) $P = .68^*$ Drug-related side effects       8 (17.8)       1 (14.3) $P = .65^*$ Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ $P = .06^*$ > 1       40 (88.9)       4 (57.1)         1       5 (11.1)       3 (42.9)	suicidality or psychosis			
$\begin{array}{c c} \mbox{Catatonia symptoms} & 13 (28.9) & 2 (28.6) & P = .68^* \\ \mbox{Drug-related side effects} & 8 (17.8) & 1 (14.3) & P = .65^* \\ \mbox{Secondary repetitive behavior} & 5 (11.1) & 2 (28.6) & P = .23^* \\ \mbox{Previous response to ECT or seizure} & 1 (2.2) & 0 & P = .86^* \\ \mbox{No. of ECT indications} & P = .06^* \\ \mbox{>} 1 & 40 (88.9) & 4 (57.1) \\ \hline 1 & 5 (11.1) & 3 (42.9) \end{array}$	Severe mania, psychosis, or agitation	15 (33.3)	1 (14.3)	P=.29*
$\begin{array}{c cccc} Drug-related side effects & 8 (17.8) & 1 (14.3) & P=.65^{*} \\ Secondary repetitive behavior & 5 (11.1) & 2 (28.6) & P=.23^{*} \\ Previous response to ECT or seizure & 1 (2.2) & 0 & P=.86^{*} \\ No. of ECT indications & & P=.06^{*} \\ >1 & 40 (88.9) & 4 (57.1) \\ \hline 1 & 5 (11.1) & 3 (42.9) \end{array}$	Catatonia symptoms	13 (28.9)	2 (28.6)	P=.68*
Secondary repetitive behavior       5 (11.1)       2 (28.6) $P = .23^*$ Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ > 1       40 (88.9)       4 (57.1)         1       5 (11.1)       3 (42.9)	Drug-related side effects	8 (17.8)	1 (14.3)	P=.65*
Previous response to ECT or seizure       1 (2.2)       0 $P = .86^*$ No. of ECT indications $P = .06^*$ > 1       40 (88.9)       4 (57.1)         1       5 (11.1)       3 (42.9)	Secondary repetitive behavior	5 (11.1)	2 (28.6)	P=.23*
No. of ECT indications         P=.06*           >1         40 (88.9)         4 (57.1)           1         5 (11.1)         3 (42.9)	Previous response to ECT or seizure	1 (2.2)	0	P=.86*
> 1 40 (88.9) 4 (57.1) 1 5 (11.1) 3 (42.9)	No. of ECT indications			P=.06*
<u> </u>	>1	40 (88.9)	4 (57.1)	
		5 (11.1)	3 (42.9)	

<sup>a</sup>Values shown as n (%) unless otherwise noted.

\*Fisher exact test.

Abbreviations: CBT = cognitive-behavioral therapy, ECT = electroconvulsive therapy, SRI = serotonin reuptake inhibitors

or tics (not explicitly involving self-injurious behaviors), 13% body dysmorphic disorder, and 2% trichotillomania (see patient features in Table 2).

The final sample included 64 patients for whom it was possible to establish a positive response to ECT, which was observed in 73.4% of the total cases, including 44.0% of the body dysmorphic disorder, 74.1% of the tic disorder, and 85.7% of the self-injurious behavior patients. Nevertheless, only 12 (17.4%) cases had formal assessments of the severity of their OCRDs before and after the treatment. Further, follow-up was described in only 27 individuals (39.1%), ranging from 1 week to 48 months. Although relapse was noted in more than half (55.6%) of individuals who were followed up, positive responses were reported in all individuals (n = 12) who received a second course of ECT.

We could establish the stimulus electrode placement in 45 OCRD cases: ECT was bilateral in 36 (80.0%), unilateral in 6 (13.3%), and unilateral followed by bilateral or vice versa in 3 (6.7%) patients. In 24 cases, mode of ECT placement was unclear. The mean (SD) number of ECT sessions administered in the first course of ECT, based on a total

of 51 available cases, was 11.27 (5.73). Frequency of ECT administration was established in 21 patients. The mean (SD) number of ECT sessions per week was initially 2.95 (0.50). There was a reduction in the frequency of ECT in the first treatment course in 20 (37.0%) cases.

Individuals who did and did not respond to ECT were compared and contrasted (Table 3). Groups did not differ in terms of age, sex, age at OCRD onset, course and pattern of comorbidity, indication for ECT, electrode placement, total number of sessions, and frequency of administration. However, there was a trend for higher frequency of selfinjurious behaviors between responders (P = .08). Articles describing ECT responders were more likely to report previous unsuccessful treatment with antipsychotics (P < .001), SRIs (P = .007), and CBT (P = .05). However, after the Benjamini-Hochberg procedure was applied, only differences in antipsychotic and SRI exposure remained significant. Publication bias was investigated by comparing the positive responses described in case reports (68.1%) to those reported in case series (n > 1) (31.9%), but no significant differences emerged ( $\chi^2_1 = 2.3, P = .12$ ).

#### dos Santos-Ribeiro et al **It is illegal to post this copyrighted PDF on any website.** Bupport the use of ECT in an atypical population that is not

In this systematic review, we were unable to find any RCT on the use of ECT in OCRD patients. As such, it is not possible to provide unequivocal evidence of the efficacy of ECT in this group of individuals. Although a positive response was reported in 73.4% of published patients, all records were case reports and series, whose ability to provide reliable evidence of treatment benefits are generally undermined by a high probability of selection, information, and, frequently, publication biases.<sup>102,103</sup> Accordingly, case reports and series lie at the penultimate level in the hierarchy of evidence of treatment benefits, being only superior to mechanistic reasoning.<sup>104</sup> However, reviews of such cases have been reported as alternative "clinician-friendly methods" to test hypothetical therapeutic models.<sup>103,105</sup> This theory is consistent with the finding that up to 22% of cases published in a high-impact journal have been followed by trials after a period of sometimes only 5 years.<sup>106</sup> Thus, the available evidence does suggest that an RCT of ECT in OCRDs may be warranted, particularly in more severe tic disorder and self-injurious behavior cases.

We also found that less than 40% of all OCRD patients treated with ECT were evaluated after administration of the index course of ECT (ie, followed up). However, when these follow-up assessments were reported, relapses were noted in more than half of individuals. Similarly high rates of post-ECT relapse in the first year have been described in depression (51% in 1 year)<sup>107</sup> and schizophrenia (42.7 to 63.6%).<sup>108,109</sup> Fortunately, all OCRD individuals who relapsed responded favorably to a repeated course of ECT. High relapse rates coupled with prompt remission after treatment resumption are consistent with an ability of ECT to keep symptoms suppressed until spontaneous remission occurs.<sup>110</sup> As many OCRDs follow a chronic course, future studies should examine the role of maintenance ECT in this specific population. Maintenance ECT has been used safely for up to 12 years<sup>111</sup> and proved effective in depression,<sup>112,113</sup> bipolar disorder,<sup>114</sup> and schizophrenia.<sup>115</sup> However, the cost of sessions and concerns of patients and families about the side effects of prolonged ECT may contribute to withdrawal and discontinuation of maintenance ECT.<sup>116</sup>

Although ECT responders and nonresponders did not differ in most parameters, there was a trend for higher frequency of self-injurious behaviors and greater exposure to unsuccessful trials with antipsychotics and SRIs among the former group. These findings are in sharp contrast to our previous review,<sup>8</sup> in which ECT responders were less likely to have been treated with SRIs or CBT in the past. It is difficult to speculate on the significance of these findings, but they suggest that either (1) ECT is more effective as an augmentation strategy for patients under existing anti-OCRD treatments compared to patients who are not using antipsychotics or SRIs or (2) authors who describe response of OCRD patients to ECT are more likely to emphasize or report previous resistance to antipsychotics or SRIs. We tend to favor the second interpretation, as it may be difficult to usually seen as responsive to ECT.

Our study has a number of limitations. First, it partly consisted of an analysis of individual cases of OCRDs treated with ECT as reported in the literature, which are subject to missing data. Second, reviewed cases comprised a heterogeneous sample with self-injurious behaviors (69.2%), tic disorders and Tourette's disorder (35.4%), body dysmorphic disorder (13%), and trichotillomania (2%). However, our study should be considered consistent with contemporary models (eg, Research Domain Criteria) suggesting that OCRDs may respond to treatments that cut across traditional diagnostic boundaries. Third, only 12 (17.4%) cases had formal assessments of severity, and no transdiagnostic measure of efficacy was available, so we had to rely on authors' descriptions of treatment response to report efficacy and compare responders to nonresponders. Finally, although it would be interesting to investigate whether risk factors for relapse reflect features related to the underlying OCRD or the initial ECT course,<sup>117</sup> low numbers precluded us from comparing patients that did versus did not relapse.

#### CONCLUSIONS

The finding that more than 70% of the reviewed cases (including body dysmorphic disorder, tic disorders, autism, and self-injurious behaviors) have shown some response to ECT should not be considered unequivocal evidence of its efficacy in OCRDs. The primary limitation to forming any firm conclusions is that there is currently a total lack of RCTs on the efficacy of ECT in different OCRDs. An additional limitation is that there is an absence of valid transdiagnostic tools for the assessment of treatment response across different OCRDs. Nonetheless, the available evidence does suggest that an RCT of ECT in OCRDs may be warranted, particularly in more severe tic disorders and self-injurious behavior cases in which maintenance ECT may be advisable due to the high relapse rate. The reported association between OCRD response to ECT and previous treatment with antipsychotics and antidepressants suggests that either (1) ECT is more effective as an augmentation strategy for existing anti-OCRD treatments or (2) authors who describe response to ECT are more likely to emphasize or report previous resistance to antipsychotics or antidepressants.

Submitted: June 16, 2018; accepted August 7, 2018. Published online: October 18, 2018.

Potential conflicts of interest: None.

**Funding/support:** This work was supported by Conselho Nacional de Desenvolvimento Científico e Tecnológico (Dr Fontenelle, grant no. 308237/2014-5), Fundação Carlos Chagas Filho de Amparo à Pesquisa do Estado do Rio de Janeiro (Dr Fontenelle), the D'Or Institute of Research and Education (Dr Fontenelle), the David Winston Turner Endowment Fund (Drs Fontenelle and Yücel); and the National Health and Medical Research Council of Australia (Dr Yücel, grant no. APP1117188).

**Role of the sponsor:** The supporters had no role in the design, analysis, interpretation, or publication of this study.

**Previous presentation:** This study was presented as an abstract at the 31st European Congress of Neuropsychopharmacology; October 6–9 2018; Barcelona, Spain.

Supplementary material: See accompanying pages.

#### ECT in Obsessive-Compulsive-Related Disorders **It is illegal to post this copyrighted** T. Cavanna AE, Robertson MM, Critchley HD. **PDF** 00, 24(3-4), 141-145. **PDF** 1995,64(3-4), 141-145.

- American Psychiatric Association. *Diagnostic* and Statistical Manual for Mental Disorders. Fifth Edition. Washington, DC: American Psychiatric Association; 2013.
- Ruscio AM, Stein DJ, Chiu WT, et al. The epidemiology of obsessive-compulsive disorder in the National Comorbidity Survey Replication. *Mol Psychiatry*. 2010;15(1):53–63.
- Sookman D, Fineberg NA; Accreditation Task Force of The Canadian Institute for Obsessive Compulsive Disorders. Specialized psychological and pharmacological treatments for obsessive-compulsive disorder throughout the lifespan: a special series by the Accreditation Task Force (ATF) of The Canadian Institute for Obsessive Compulsive Disorders (CIOCD, www.ciocd.ca). Psychiatry Res. 2015;227(1):74–77.
- Simpson HB, Foa EB, Liebowitz MR, et al. Cognitive-behavioral therapy vs risperidone for augmenting serotonin reuptake inhibitors in obsessive-compulsive disorder: a randomized clinical trial. JAMA Psychiatry. 2013;70(11):1190–1199.
- Grøtte T, Hansen B, Haseth S, et al. Three-week inpatient treatment of obsessive-compulsive disorder: a 6-month follow-up study. Front Psychol. 2018;9:620.
- Alonso P, Cuadras D, Gabriëls L, et al. Deep brain stimulation for obsessive-compulsive disorder: a meta-analysis of treatment outcome and predictors of response. *PLoS One*. 2015;10(7):e0133591.
- Miguel EC, Lopes AC, McLaughlin NCR, et al. Evolution of gamma knife capsulotomy for intractable obsessive-compulsive disorder [published online ahead of print May 9, 2018]. *Mol Psychiatry.*
- Lins-Martins NM, Yücel M, Tovar-Moll F, et al. Electroconvulsive therapy in obsessivecompulsive disorder: a chart review and evaluation of its potential therapeutic effects. J Neuropsychiatry Clin Neurosci. 2015;27(1):65–68.
- Fontenelle LF, Coutinho ES, Lins-Martins NM, et al. Electroconvulsive therapy for obsessivecompulsive disorder: a systematic review. J Clin Psychiatry. 2015;76(7):949–957.
- Phillips KA, Stein DJ, Rauch SL, et al. Should an obsessive-compulsive spectrum grouping of disorders be included in DSM-V? Depress Anxiety. 2010;27(6):528–555.
- Taylor S. Early versus late onset obsessivecompulsive disorder: evidence for distinct subtypes. *Clin Psychol Rev.* 2011;31(7):1083–1100.
- de Vries FE, Cath DC, Hoogendoorn AW, et al. Tic-related versus tic-free obsessivecompulsive disorder: clinical picture and 2-year natural course. J Clin Psychiatry. 2016;77(10):e1240–e1247.
- Harris KM, Mahone EM, Singer HS. Nonautistic motor stereotypies: clinical features and longitudinal follow-up. *Pediatr Neurol*. 2008;38(4):267–272.
- Phillips KA, Pinto A, Hart AS, et al. A comparison of insight in body dysmorphic disorder and obsessive-compulsive disorder. *J Psychiatr Res.* 2012;46(10):1293–1299.
- Torres AR, Fontenelle LF, Ferrão YA, et al. Clinical features of obsessive-compulsive disorder with hoarding symptoms: a multicenter study. *J Psychiatr Res.* 2012;46(6):724–732.
- Phillips KA, Didie ER, Menard W. Clinical features and correlates of major depressive disorder in individuals with body dysmorphic disorder. J Affect Disord. 2007;97(1–3):129–135.

Cavanna AE, Robertson MM, Critchley HD: Catatonic signs in Gilles de la Tourette syndrome. *Cogn Behav Neurol*. 2008;21(1):34–37.

- Dhossche DM, Reti IM, Shettar SM, et al. Tics as signs of catatonia: electroconvulsive therapy response in 2 men. *J ECT*. 2010;26(4):266–269.
- DeJong H, Bunton P, Hare DJ. A systematic review of interventions used to treat catatonic symptoms in people with autistic spectrum disorders. J Autism Dev Disord. 2014;44(9):2127–2136.
- Dhossche DM. Decalogue of catatonia in autism spectrum disorders. *Front Psychiatry*. 2014;5:157.
- Mazzone L, Postorino V, Valeri G, et al. Catatonia in patients with autism: prevalence and management. CNS Drugs. 2014;28(3):205–215.
- Wachtel LE, Shorter E. Self-injurious behaviour in children: A treatable catatonic syndrome. *Aust N Z J Psychiatry*. 2013;47(12):1113–1115.
- O'Sullivan RL, Phillips KA, Keuthen NJ, et al. Near-fatal skin picking from delusional body dysmorphic disorder responsive to fluvoxamine. *Psychosomatics*. 1999;40(1):79–81.
- 24. Darke S, Duflou J. Characteristics, circumstances and pathology of sudden or unnatural deaths of cases with evidence of pathological hoarding. *J Forensic Leg Med*. 2017;45:36–40.
- Kırpınar I, Kocacenk T, Koçer E, et al. Recurrent trichobezoar due to trichophagia: a case report. *Gen Hosp Psychiatry*. 2013;35(4):439–441.
- Weintraub E, Robinson C, Newmeyer M. Catastrophic medical complication in psychogenic excoriation. South Med J. 2000;93(11):1099–1101.
- 27. Cheung MY, Shahed J, Jankovic J. Malignant Tourette syndrome. *Mov Disord*. 2007;22(12):1743–1750.
- Wachtel LE, Griffin M, Reti IM. Electroconvulsive therapy in a man with autism experiencing severe depression, catatonia, and self-injury. *J ECT*. 2010;26(1):70–73.
- Kalinowsky LB, Hippius H. Pharmacological, Convulsive, and Other Somatic Treatments in Psychiatry. New York, NY: Grune & Stratton; 1969.
- Shamseer L, Moher D, Clarke M, et al; PRISMA-P Group. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015: elaboration and explanation. *BMJ*. 2015;350:g7647.
- Phillips KA, Hollander E, Rasmussen SA, et al. A severity rating scale for body dysmorphic disorder: development, reliability, and validity of a modified version of the Yale-Brown Obsessive Compulsive Scale. *Psychopharmacol Bull.* 1997;33(1):17–22.
- Arnold LM, Mutasim DF, Dwight MM, et al. An open clinical trial of fluvoxamine treatment of psychogenic excoriation. J Clin Psychopharmacol. 1999;19(1):15–18.
- Frost RO, Steketee G, Grisham J. Measurement of compulsive hoarding: Saving Inventory-Revised. *Behav Res Ther*. 2004;42(10):1163–1182.
- O'Sullivan RL, Keuthen NJ, Hayday CF, et al. The Massachusetts General Hospital (MGH) Hairpulling Scale: 2. reliability and validity. *Psychother Psychosom*. 1995;64(3–4):146–148.
- Keuthen NJ, O'Sullivan RL, Ricciardi JN, et al. The Massachusetts General Hospital (MGH) Hairpulling Scale: 1. development and factor analyses. *Psychother Psychosom*.

- Leckman JF, Riddle MA, Hardin MT, et al. The Yale Global Tic Severity Scale: initial testing of a clinician-rated scale of tic severity. J Am Acad Child Adolesc Psychiatry. 1989;28(4):566–573.
- Scahill L, McDougle CJ, Williams SK, et al; Research Units on Pediatric Psychopharmacology Autism Network. Children's Yale-Brown Obsessive Compulsive Scale modified for pervasive developmental disorders. J Am Acad Child Adolesc Psychiatry. 2006;45(9):1114–1123.
- Grant JE, Kim SW, Odlaug BL. A double-blind, placebo-controlled study of the opiate antagonist, naltrexone, in the treatment of kleptomania. *Biol Psychiatry*. 2009;65(7):600–606.
- Coffey CE. The Clinical Science of Electroconvulsive Therapy. Washington, DC: American Psychiatric Press; 1993.
- Shorter E, Healy D. Shock Therapy: A History of Electroconvulsive Treatment in Mental Illness. New Brunswick, NJ: Rutgers University Press; 2007.
- Mankad MV, Beyer JL, Weiner RD, et al. Clinical Manual of Electroconvulsive Therapy. Arlington, VA: American Psychiatric Publishing, Incorporated; 2010.
- McDonald WM, Meeks TW, McCall WV, et al. *Electroconvulsive Therapy*. Arlington, VA: American Psychiatric Publishing; 2009.
- Swartz CM. Patient selection and electroconvulsive therapy indications. *Electroconvulsive and Neuromodulation Therapies*. New York, NY: Cambridge University Press; 2009:1114–1123.
- 44. Weiner RD. The Practice of Electroconvulsive Therapy: Recommendations for Treatment, Training, and Privileging: A Task Force Report of the American Psychiatric Association. Arlington, VA: American Psychiatric Association; 2008.
- Fink M. Electroconvulsive Therapy: A Guide for Professionals and Their Patients. Cary, NC: Oxford University Press; 2009.
- Higgins JPT, Green S. Cochrane Handbook for Systematic Reviews of Interventions. Chichester, UK: Wiley; 2011.
- Benjamini Y, Hochberg Y. Controlling the false discovery rate: a practical and powerful approach to multiple testing. *J R Stat Soc B*. 1995;57(1):289–300.
- Denning S, Mehrkens JH, Müller N, et al. Therapy-refractory Tourette syndrome: beneficial outcome with globus pallidus internus deep brain stimulation. *Mov Disord*. 2008;23(9):1300–1302.
- Duits A, Ackermans L, Cath D, et al. Unfavourable outcome of deep brain stimulation in a Tourette patient with severe comorbidity. *Eur Child Adolesc Psychiatry*. 2012;21(9):529–531.
- Godart NT, Agman G, Perdereau F, et al. Naltrexone treatment of self-injurious behavior. J Am Acad Child Adolesc Psychiatry. 2000;39(9):1076–1078.
- Houeto JL, Karachi C, Mallet L, et al. Tourette's syndrome and deep brain stimulation. *J Neurol Neurosurg Psychiatry*. 2005;76(7):992–995.
- Sperling W, Reulbach U, Maihöfner C, et al. Vagus nerve stimulation in a patient with Gilles de la Tourette syndrome and major depression. *Pharmacopsychiatry*. 2008;41(3):117–118.
- Dehning S, Feddersen B, Cerovecki A, et al. Globus pallidus internus-deep brain stimulation in Tourette's syndrome: can clinical symptoms predict response? *Mov*

# It is illegal to post this copyrighted PDF, on any website.

- Corbella T, Rossi L. Dysmorphophobia: its clinical and nosographical aspects. Acta Neurol Psychiatr Belg. 1967;67(9):691–700.
- Araneta E, Magen J, Musci MN Jr, et al. Gilles de la Tourette's syndrome symptom onset at age 35. *Child Psychiatry Hum Dev.* 1975;5(4):224–230.
- Bates WJ, Smeltzer DJ. Electroconvulsive treatment of psychotic self-injurious behavior in a patient with severe mental retardation. *Am J Psychiatry.* 1982;139(10):1355–1356.
- Guttmacher LB, Cretella H. Electroconvulsive therapy in one child and three adolescents. *J Clin Psychiatry*. 1988;49(1):20–23.
- Swerdlow NR, Gierz M, Berkowitz A, et al. Electroconvulsive therapy in a patient with severe tic and major depressive episode. *J Clin Psychiatry*. 1990;51(1):34–35.
- Phillips KA, McElroy SL, Keck PE Jr, et al. Body dysmorphic disorder: 30 cases of imagined ugliness. *Am J Psychiatry*. 1993;150(2):302–308.
- Carroll BJ, Yendrek R, Degroot C, et al. Response of major depression with psychosis and body dysmorphic disorder to ECT. Am J Psychiatry. 1994;151(2):288–289.
- Cizadlo BC, Wheaton A. Case study: ECT treatment of a young girl with catatonia. J Am Acad Child Adolesc Psychiatry. 1995;34(3):332–335.
- Rapoport M, Feder V, Sandor P. Response of major depression and Tourette's syndrome to ECT: a case report. *Psychosom Med.* 1998;60(4):528–529.
- Dean CE. Severe self-injurious behavior associated with treatment-resistant schizophrenia: treatment with maintenance electroconvulsive therapy. *J ECT*. 2000;16(3):302–308.
- Myers W, Nguyen M. Modified multiplemonitored electroconvulsive therapy. J Am Acad Child Adolesc Psychiatry. 2002;41(7):756–758.
- Friedlander RI, Solomons K. ECT: use in individuals with mental retardation. *JECT*. 2002;18(1):38–42.
- Trivedi HK, Mendelowitz AJ, Fink M. Gilles de la Tourette form of catatonia: response to ECT. *J ECT*. 2003;19(2):115–117.
- Kessler RJ. Electroconvulsive therapy for affective disorders in persons with mental retardation. *Psychiatr Q*. 2004;75(1):99–104.
- 68. Strassnig M, Riedel M, Muller N. Electroconvulsive therapy in a patient with Tourette's syndrome and co-morbid Obsessive Compulsive Disorder. The world journal of biological psychiatry : the official journal of the World Federation of Societies of Biological Psychiatry. 2004;5(3):164-166.
- 69. Kelly VC, Chan YC. Oedipism thwarted with electroconvulsive therapy. *J ECT*. 2004;20(4):273–274.
- Karadenizli D, Dilbaz N, Bayam G. Gilles de la Tourette syndrome: response to electroconvulsive therapy. *J ECT*. 2005;21(4):246–248.
- Ghaziuddin M, Quinlan P, Ghaziuddin N. Catatonia in autism: a distinct subtype? J Intellect Disabil Res. 2005;49(Pt 1):102–105.
- Morais SL, Derenusson GN, Pinto JP, et al. Neurobiological substrates of electroconvulsive therapy for Tourette syndrome: a Serial SISCOM study. *J ECT*. 2007;23(4):278–280.
- Okada T, Yamamoto H, Funabiki Y. Effectiveness of electroconvulsive therapy and donepezil treatment to severe tics in Tourette's disorder. 2007.
- 74. Fisher CE, Sporn AL, Mantovani A, et al.

Electroconvulsive therapy as an alternati deep brain stimulation for medicationrefractory tourette syndrome. *JECT*. 2008;24(1):2.

- Arora M, Praharaj SK, Prakash R. Electroconvulsive therapy for multiple major self-mutilations in bipolar psychotic depression [article in Turkish]. Turkish J Psychiatry. 2008;19(2):209–212.
- Wachtel LE, Kahng S, Dhossche DM, et al. ECT for catatonia in an autistic girl. Am J Psychiatry. 2008;165(3):329–333.
- Kakooza-Mwesige A, Wachtel LE, Dhossche DM. Catatonia in autism: implications across the life span. *Eur Child Adolesc Psychiatry*. 2008;17(6):327–335.
- Wachtel LE, Contrucci-Kuhn SA, Griffin M, et al. ECT for self-injury in an autistic boy. Eur Child Adolesc Psychiatry. 2009;18(7):458–463.
   Washel LE, Griffin M, Beitin M,
- Wachtel LE, Griffin M, Reti IM. Electroconvulsive therapy in a man with autism experiencing severe depression, catatonia, and self-injury. *J ECT*. 2010;26(1):70–73.
- Wachtel LE, Jaffe R, Kellner CH. Electroconvulsive therapy for psychotropicrefractory bipolar affective disorder and severe self-injury and aggression in an 11-year-old autistic boy. *Eur Child Adolesc Psychiatry*. 2011;20(3):147–152.
- Ghaziuddin N, Dumas S, Hodges E. Use of continuation or maintenance electroconvulsive therapy in adolescents with severe treatment-resistant depression. *J ECT*. 2011;27(2):168–174.
- Siegel M, Milligan B, Robbins D, et al. Electroconvulsive therapy in an adolescent with autism and bipolar I disorder. *J ECT*. 2012;28(4):252–255.
- Rapinesi C, Serata D, Del Casale A, et al. Effectiveness of electroconvulsive therapy in a patient with a treatment-resistant major depressive episode and comorbid body dysmorphic disorder. *JECT*. 2013;29(2):145–146.
- Consoli A, Cohen J, Bodeau N, et al.
   Electroconvulsive therapy in adolescents with intellectual disability and severe self-injurious behavior and aggression: a retrospective study. *Eur Child Adolesc Psychiatry*. 2013;22(1):55–62.
- Navinés R, Gutierrez F, Arranz B, et al. Longterm and bizarre self-injurious behavior: an approach to underlying psychological mechanisms and management. J Psychiatr Pract. 2013;19(1):65–71.
- Wachtel LE, Schuldt S, Ghaziuddin N, et al. The potential role of electroconvulsive therapy in the 'Iron Triangle' of pediatric catatonia, autism, and psychosis. *Acta Psychiatr Scand*. 2013;128(5):408–409.
- Rajashree VC, Manjiri CD, Ivan SN, et al. Gilles de la Tourette's syndrome successfully treated with electroconvulsive therapy. *Indian J Psychiatry*. 2014;56(4):407–408.
- Wachtel LE, Reti IM, Ying H. Stability of intraocular pressure after retinal reattachment surgery during electroconvulsive therapy for intractable self-injury in a 12-year-old autistic boy. *J ECT*. 2014;30(1):73–76.
- 89. Haq AU, Ghaziuddin N. Maintenance electroconvulsive therapy for aggression and self-injurious behavior in two adolescents with autism and catatonia. *J Neuropsychiatry Clin Neurosci.* 2014;26(1):64–72.
- Okazaki R, Takahashi T, Ueno K, et al. Changes in EEG complexity with electroconvulsive therapy in a patient with autism spectrum disorders: a multiscale entropy approach. *Front Hum Neurosci.* 2015;9:106.

Wachtel L, Commins E, Park M, et al. Neuroleptic malignant syndrome and delirious mania as malignant catatonia in autism: prompt relief with electroconvulsive therapy. Acta Psychiatr Scand. 2015;132(4):319–320.

- Dhossche DM, van der Steen LF, Shettar SM. Catatonia in autism spectrum disorders: review and case-report [article in Dutch]. *Tijdschr Psychiatr.* 2015;57(2):89–93.
- Mahato R<sup>5</sup>, San Gabriel MC, Longshore CT, et al. A Case of Treatment- resistant Depression and Body Dysmorphic Disorder: The Role of Electroconvulsive Therapy Revisited. Innov Clin Neurosci. 2016;13(7-8):37–40.
- Guo JN, Kothari JS, Leckman JF, et al. Successful Treatment of Tourette Syndrome With Electroconvulsive Therapy: A Case Report. *Biol Psychiatry*. 2016;79(5):e13–e14.
- Clinebell K, Valpey R, Walker T, et al. Self-Enucleation and Severe Ocular Injury in the Psychiatric Setting. *Psychosomatics*. 2016;57(1):25–30.
- Katz R, Bukanova E, Ostroff R. Procedural Consolidation During Electroconvulsive Therapy for a Patient With Severe Tourette Syndrome. JECT. 2017;33(1):e7–e8.
- Sajith SG, Liew SF, Tor PC. Response to Electroconvulsive Therapy in Patients With Autism Spectrum Disorder and Intractable Challenging Behaviors Associated With Symptoms of Catatonia. J ECT. 2017;33(1):63–67.
- Yero SA, McKinney T, Petrides G, et al. Successful use of electroconvulsive therapy in 2 cases of persistent sexual arousal syndrome and bipolar disorder. *JECT*. 2006;22(4):274–275.
- Larkin EP. Milieu and mutilation—a case for 'special' treatment? Br J Psychiatry. 1992;160(01):116–119.
- Bailine SH, Petraviciute S. Catatonia in autistic twins: role of electroconvulsive therapy. J ECT. 2007;23(1):21–22.
- 101. Thuppal M, Fink M. Electroconvulsive therapy and mental retardation. *JECT*. 1999;15(2):140–149.
- 102. García-Doval I, Albrecht J, Flohr C, et al; European Dermato-Epidemiology Network (EDEN). Optimizing case reports and case series: guidance on how to improve quality. Br J Dermatol. 2018;178(6):1257–1262.
- Vandenbroucke JP. Case reports in an evidence-based world. JR Soc Med. 1999;92(4):159–163.
- Howick J, Chalmers I, Glasziou P, et al. The 2011 Oxford CEBM Evidence Levels of Evidence (Introductory Document). Center for Evidence-Based Medicine website. http://www.cebm.net/index.aspx?o=5653. 2011. Accessed May 18, 2018.
- Charlton BG, Walston F. Individual case studies in clinical research. J Eval Clin Pract. 1998;4(2):147–155.
- Albrecht J, Meves A, Bigby M. Case reports and case series from Lancet had significant impact on medical literature. J Clin Epidemiol. 2005;58(12):1227–1232.
- 107. Kellner CH. Relapse after electroconvulsive therapy (ECT). *J ECT*. 2013;29(1):1–2.
- 108. Shibasaki C, Takebayashi M, Fujita Y, et al. Factors associated with the risk of relapse in schizophrenic patients after a response to electroconvulsive therapy: a retrospective study. *Neuropsychiatr Dis Treat*. 2015;11:67–73.
- 109. Suzuki K, Awata S, Matsuoka H. One-year outcome after response to ECT in middleaged and elderly patients with intractable

#### ECT in Obsessive-Compulsive-Related Disorders

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2004;20(2):99-106.

- 110. Fox HA. Continuation and maintenance electroconvulsive therapy-a conceptual framework? JECT. 2015;31(2):e27-e28.
- 111. Elias A, Chathanchirayil SJ, Bhat R, et al. Maintenance electroconvulsive therapy up to 12 years. J Affect Disord. 2014;156:228-231.
- 112. Elias A, Phutane VH, Clarke S, et al. Electroconvulsive therapy in the continuation and maintenance treatment of depression: Systematic review and meta-analyses. Aust N

113. Gill SP, Kellner CH. Clinical practice

- recommendations for continuation and maintenance electroconvulsive therapy for depression: outcomes from a review of the evidence and a consensus workshop held in Australia in May 2017 [published online ahead of print February 7, 2018]. JECT.
- 114. Santos Pina L, Bouckaert F, Obbels J, et al. Maintenance electroconvulsive therapy in severe bipolar disorder: a retrospective chart review. JECT. 2016;32(1):23-28.

Ward HB, Szabo ST, Rakesh G. Maintenance ECT in schizophrenia: a systematic review. Psychiatry Res. 2018;264:131-142.

- 116. Rodriguez-Jimenez R, Bagney A, Torio I, et al. Maintenance electroconvulsive therapy: costeffectiveness and patient/family Satisfaction. JECT. 2015;31(4):279.
- Petrides G, Tobias KG, Kellner CH, et al. 117. Continuation and maintenance electroconvulsive therapy for mood disorders: review of the literature. Neuropsychobiology. 2011;64(3):129-140.

Supplementary material follows this article.

## THE PRIMARY CARE COMPANION FOR CNS DISORDERS

### **Supplementary Material**

Article Title: A Systematic Review of the Utility of Electroconvulsive Therapy in Broadly Defined Obsessive-Compulsive–Related Disorders

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DOI Number: https://doi.org/10.4088/PCC.18r02342

### List of Supplementary Material for the article

1. Supplementary Table 1. List of Studies Included in the Systematic Review

#### **Disclaimer**

This Supplementary Material has been provided by the author(s) as an enhancement to the published article. It has been approved by peer review; however, it has undergone neither editing nor formatting by in-house editorial staff. The material is presented in the manner supplied by the author.

Study first author and year	Country	Design	ECT patients, N	Main indication for ECT
Corbella, 1967 <sup>1</sup>	France	Case series	2	Unclear
Araneta, 1975 <sup>2</sup>	USA	Case report	1	Treatment resistance
Bates, 1982 <sup>3</sup>	USA	Case report	1	More than one
Guttmacher, 1988 <sup>4</sup>	USA	Case series	1	More than one
Swerdlow, 1990 <sup>5</sup>	USA	Case report	1	More than one
Larkin, 1992 <sup>6</sup>	UK	Case report	1	Unclear
Phillips, 1993 <sup>7</sup>	USA	Case series	4	Unclear
Carroll, 1994 <sup>8</sup>	USA	Case report	1	Severe Major depression
Cizadlo, 1995 <sup>9</sup>	USA	Case report	1	More than one
Rapoport, 1998 <sup>10</sup>	Canada	Case report	1	Severe Major depression
Thuppal, 1999 <sup>11</sup>	USA	Case series	1	Severe Major depression
Dean, 2000 <sup>12</sup>	USA	Case report	1	More than one
Godart, 2000 <sup>13</sup>	France	Case report	1	Unclear
Myers, 2002 <sup>14</sup>	USA	Case report	1	More than one
Friedlander, 2002 <sup>15</sup>	Canada	Case series	1	More than one
Trivedi, 2003 <sup>16</sup>	USA	Case report	1	More than one
Kessler, 2004 <sup>17</sup>	USA	Case series	1	Severe Major depression
Strassning, 2004 <sup>18</sup>	Germany	Case report	1	Previous response to ECT or convulsion
Kelly, 2004 <sup>19</sup>	USA	Case report	1	More than one
Karadenizeli, 2005 <sup>20</sup>	Turkey	Case report	1	More than one
Houeto, 2005 <sup>21</sup>	France	Case report	1	Unclear
Ghaziuddin, 2005 <sup>22</sup>	USA	Case report	1	More than one
Yero, 2006 <sup>23</sup>	USA	Case series	1	Others
Morais, 2007 <sup>24</sup>	Brazil	Case report	1	More than one
Okada, 2007 <sup>25</sup>	Japan	Case report	1	More than one
Bailine, 2007 <sup>26</sup>	USA	Case series	1	More than one
Dehning, 2008 <sup>27</sup>	Germany	Case report	1	More than one
Sperling, 2008 <sup>28</sup>	Germany	Case report	1	Treatment resistance
Fisher, 2008 <sup>29</sup>	USA	Case report	1	More than one
Arora, 2008 <sup>30</sup>	India	Case report	1	More than one
Wachtel, 2008 <sup>31</sup>	USA	Case report	1	More than one
Kakooza-Mwesige, 2008 <sup>32</sup>	-	Case report	1	More than one
Wachtel, 2009 <sup>33</sup>	USA	Case report	1	More than one
Fink, 2009 <sup>34</sup>	USA	Case report	1	More than one
Dhossche, 2010 <sup>35</sup>	USA	Case series	2	More than one
Watchel, 2010 <sup>36</sup>	USA	Case report	1	More than one
Dehning, 2011 <sup>37</sup>	Germany	Case series	4	Unclear
Dehning, 2011 <sup>38</sup>	Germany	Case report	1	More than one
Watchel, 2011 <sup>39</sup>	USA	Case report	1	More than one

Supplementary Table 1. List of Studies Included in the Systematic Review

Ghaziuddin, 2011 <sup>40</sup>	USA	Case series	2	More than one
Duits, 2012 <sup>41</sup>	Netherlands	Case report	1	More than one
Siegel, 2012 <sup>42</sup>	USA	Case report	1	More than one
Rapinesi, 2013 <sup>43</sup>	Italy	Case report	1	Treatment resistance
Consoli, 2013 <sup>44</sup>	France	Case series	4	More than one
Navinés, 201345	Spain	Case report	1	Unclear
Watchel, 2013 <sup>46</sup>	USA	Case report	1	More than one
Rajashree, 2014 <sup>47</sup>	India	Case report	1	More than one
Watchel, 2014 <sup>48</sup>	USA	Case report	1	More than one
Haq, 2014 <sup>49</sup>	USA	Case series	1	More than one
Okazaki, 2015 <sup>50</sup>	Japan	Case report	1	Severe Catatonia
Watchel, 2015 <sup>51</sup>	USA	Case report	1	More than one
Dhossche, 2015 <sup>52</sup>	USA	Case report	1	More than one
Mahato, 2016 <sup>53</sup>	USA	Case report	1	More than one
Guo, 2016 <sup>54</sup>	USA	Case report	1	More than one
Clinebell, 2016 <sup>55</sup>	USA	Case series	1	More than one
Katz, 2017 <sup>56</sup>	USA	Case report	1	More than one
Sajith, 2017 <sup>57</sup>	Singapore	Case series	2	More than one

#### References

- 1. Corbella T, Rossi L. [Dysmorphophobia. Its clinical and nosographical aspects]. *Acta neurologica et psychiatrica Belgica*. 1967;67(9):691-700.
- 2. Araneta E, Magen J, Musci MN, Jr., Singer P, Vann CR. Gilles de la Tourette's syndrome symptom onset at age 35. *Child Psychiatry Hum Dev.* 1975;5(4):224-230.
- 3. Bates WJ, Smeltzer DJ. Electroconvulsive treatment of psychotic self-injurious behavior in a patient with severe mental retardation. *The American journal of psychiatry*. 1982;139(10):1355-1356.
- 4. Guttmacher LB, Cretella H. Electroconvulsive therapy in one child and three adolescents. *J Clin Psychiatry*. 1988;49(1):20-23.
- 5. Swerdlow NR, Gierz M, Berkowitz A, Nemiroff R, Lohr J. Electroconvulsive therapy in a patient with severe tic and major depressive episode. *J Clin Psychiatry*. 1990;51(1):34-35.
- 6. Larkin EP. Milieu and mutilation--a case for 'special' treatment? *The British journal of psychiatry : the journal of mental science*. 1992;160:116-119.
- Phillips KA, McElroy SL, Keck PE, Jr., Pope HG, Jr., Hudson JI. Body dysmorphic disorder: 30 cases of imagined ugliness. *The American journal of psychiatry*. 1993;150(2):302-308.
- 8. Carroll BJ, Yendrek R, Degroot C, Fanin H. Response of major depression with psychosis and body dysmorphic disorder to ECT. *The American journal of psychiatry*. 1994;151(2):288-289.
- 9. Cizadlo BC, Wheaton A. Case study: ECT treatment of a young girl with catatonia. *Journal of the American Academy of Child and Adolescent Psychiatry*. 1995;34(3):332-335.
- 10. Rapoport M, Feder V, Sandor P. Response of major depression and Tourette's syndrome to ECT: a case report. *Psychosomatic medicine*. 1998;60(4):528-529.

- 11. Thuppal M, Fink M. Electroconvulsive therapy and mental retardation. *The journal of ECT*. 1999;15(2):140-149.
- 12. Dean CE. Severe self-injurious behavior associated with treatment-resistant schizophrenia: treatment with maintenance electroconvulsive therapy. *The journal of ECT*. 2000;16(3):302-308.
- 13. Godart NT, Agman G, Perdereau F, Jeanmet P. Naltrexone treatment of self-injurious behavior. *J Am Acad Child Adolesc Psychiatry*. 2000;39(9):1076-1078.
- Myers W, Nguyen M. Modified multiple-monitored electroconvulsive therapy. Journal of the American Academy of Child and Adolescent Psychiatry. 2002;41(7):756-758.
- 15. Friedlander RI, Solomons K. ECT: use in individuals with mental retardation. *The journal of ECT*. 2002;18(1):38-42.
- 16. Trivedi HK, Mendelowitz AJ, Fink M. Gilles de la Tourette form of catatonia: response to ECT. *The journal of ECT*. 2003;19(2):115-117.
- 17. Kessler RJ. Electroconvulsive therapy for affective disorders in persons with mental retardation. *The Psychiatric quarterly*. 2004;75(1):99-104.
- 18. Strassnig M, Riedel M, Muller N. Electroconvulsive therapy in a patient with Tourette's syndrome and co-morbid Obsessive Compulsive Disorder. *The world journal of biological psychiatry : the official journal of the World Federation of Societies of Biological Psychiatry*. 2004;5(3):164-166.
- 19. Kelly VC, Chan YC. Oedipism thwarted with electroconvulsive therapy. *The journal* of ECT. 2004;20(4):273-274.
- 20. Karadenizli D, Dilbaz N, Bayam G. Gilles de la Tourette syndrome: response to electroconvulsive therapy. *The journal of ECT*. 2005;21(4):246-248.
- 21. Houeto JL, Karachi C, Mallet L, et al. Tourette's syndrome and deep brain stimulation. *Journal of neurology, neurosurgery, and psychiatry.* 2005;76(7):992-995.
- 22. Ghaziuddin M, Quinlan P, Ghaziuddin N. Catatonia in autism: a distinct subtype? *Journal of intellectual disability research : JIDR*. 2005;49(Pt 1):102-105.
- 23. Yero SA, McKinney T, Petrides G, Goldstein I, Kellner CH. Successful use of electroconvulsive therapy in 2 cases of persistent sexual arousal syndrome and bipolar disorder. *The journal of ECT*. 2006;22(4):274-275.
- 24. Morais SL, Derenusson GN, Pinto JP, et al. Neurobiological substrates of electroconvulsive therapy for Tourette syndrome: a Serial SISCOM study. *The journal of ECT*. 2007;23(4):278-280.
- 25. Okada T, Yamamoto H, Funabiki Y. *Effectiveness of electroconvulsive therapy and donepezil treatment to severe tics in Tourette's disorder*. 2007.
- 26. Bailine SH, Petraviciute S. Catatonia in autistic twins: role of electroconvulsive therapy. *The journal of ECT*. 2007;23(1):21-22.
- 27. Dehning S, Mehrkens JH, Muller N, Botzel K. Therapy-refractory Tourette syndrome: beneficial outcome with globus pallidus internus deep brain stimulation. *Movement disorders : official journal of the Movement Disorder Society*. 2008;23(9):1300-1302.
- 28. Sperling W, Reulbach U, Maihofner C, Kornhuber J, Bleich S. Vagus nerve stimulation in a patient with Gilles de la Tourette syndrome and major depression. *Pharmacopsychiatry*. 2008;41(3):117-118.
- 29. Fisher CE, Sporn AL, Mantovani A, Leckman JF, Lisanby SH. Electroconvulsive Therapy as an Alternative to Deep Brain Stimulation for Medication-Refractory Tourette Syndrome. *The journal of ECT*. 2008;24(1):2.
- 30. Arora M, Praharaj SK, Prakash R. [Electroconvulsive therapy for multiple major selfmutilations in bipolar psychotic depression]. *Turk psikiyatri dergisi = Turkish journal of psychiatry*. 2008;19(2):209-212.

- 31. Wachtel LE, Kahng S, Dhossche DM, Cascella N, Reti IM. ECT for catatonia in an autistic girl. *The American journal of psychiatry*. 2008;165(3):329-333.
- 32. Kakooza-Mwesige A, Wachtel LE, Dhossche DM. Catatonia in autism: implications across the life span. *Eur Child Adolesc Psychiatry*. 2008;17(6):327-335.
- Wachtel LE, Contrucci-Kuhn SA, Griffin M, Thompson A, Dhossche DM, Reti IM. ECT for self-injury in an autistic boy. *Eur Child Adolesc Psychiatry*. 2009;18(7):458-463.
- 34. Fink M. *Electroconvulsive Therapy: A Guide for Professionals and their patients* 2009.
- 35. Dhossche DM, Reti IM, Shettar SM, Wachtel LE. Tics as signs of catatonia: electroconvulsive therapy response in 2 men. *The journal of ECT*. 2010;26(4):266-269.
- 36. Wachtel LE, Griffin M, Reti IM. Electroconvulsive therapy in a man with autism experiencing severe depression, catatonia, and self-injury. *The journal of ECT*. 2010;26(1):70-73.
- 37. Dehning S, Feddersen B, Cerovecki A, Botzel K, Muller N, Mehrkens JH. Globus pallidus internus-deep brain stimulation in Tourette's syndrome: can clinical symptoms predict response? *Movement disorders : official journal of the Movement Disorder Society*. 2011;26(13):2440-2441.
- 38. Dehning S, Feddersen B, Mehrkens JH, Muller N. Long-term results of electroconvulsive therapy in severe Gilles de la Tourette syndrome. *The journal of ECT*. 2011;27(2):145-147.
- 39. Wachtel LE, Jaffe R, Kellner CH. Electroconvulsive therapy for psychotropicrefractory bipolar affective disorder and severe self-injury and aggression in an 11year-old autistic boy. *Eur Child Adolesc Psychiatry*. 2011;20(3):147-152.
- 40. Ghaziuddin N, Dumas S, Hodges E. Use of continuation or maintenance electroconvulsive therapy in adolescents with severe treatment-resistant depression. *The journal of ECT*. 2011;27(2):168-174.
- 41. Duits A, Ackermans L, Cath D, Visser-Vandewalle V. Unfavourable outcome of deep brain stimulation in a Tourette patient with severe comorbidity. *Eur Child Adolesc Psychiatry*. 2012;21(9):529-531.
- 42. Siegel M, Milligan B, Robbins D, Prentice G. Electroconvulsive therapy in an adolescent with autism and bipolar I disorder. *The journal of ECT*. 2012;28(4):252-255.
- 43. Rapinesi C, Serata D, Del Casale A, et al. Effectiveness of electroconvulsive therapy in a patient with a treatment-resistant major depressive episode and comorbid body dysmorphic disorder. *The journal of ECT.* 2013;29(2):145-146.
- 44. Consoli A, Cohen J, Bodeau N, Guinchat V, Wachtel L, Cohen D. Electroconvulsive therapy in adolescents with intellectual disability and severe self-injurious behavior and aggression: a retrospective study. *Eur Child Adolesc Psychiatry*. 2013;22(1):55-62.
- 45. Navines R, Gutierrez F, Arranz B, et al. Long-term and bizarre self-injurious behavior: an approach to underlying psychological mechanisms and management. *J Psychiatr Pract.* 2013;19(1):65-71.
- 46. Wachtel LE, Schuldt S, Ghaziuddin N, Shorter E. The potential role of electroconvulsive therapy in the 'Iron Triangle' of pediatric catatonia, autism, and psychosis. *Acta psychiatrica Scandinavica*. 2013;128(5):408-409.
- 47. Rajashree VC, Manjiri CD, Ivan SN, Alka VP. Gilles de la Tourette's syndrome successfully treated with electroconvulsive therapy. *Indian journal of psychiatry*. 2014;56(4):407-408.

- 48. Wachtel LE, Reti IM, Ying H. Stability of intraocular pressure after retinal reattachment surgery during electroconvulsive therapy for intractable self-injury in a 12-year-old autistic boy. *The journal of ECT*. 2014;30(1):73-76.
- 49. Haq AU, Ghaziuddin N. Maintenance electroconvulsive therapy for aggression and self-injurious behavior in two adolescents with autism and catatonia. *The Journal of neuropsychiatry and clinical neurosciences*. 2014;26(1):64-72.
- 50. Okazaki R, Takahashi T, Ueno K, et al. Changes in EEG complexity with electroconvulsive therapy in a patient with autism spectrum disorders: a multiscale entropy approach. *Frontiers in human neuroscience*. 2015;9:106.
- 51. Wachtel L, Commins E, Park M, Rolider N, Stephens R, Reti I. Neuroleptic malignant syndrome and delirious mania as malignant catatonia in autism: prompt relief with electroconvulsive therapy. *Acta psychiatrica Scandinavica*. 2015;132(4):319-320.
- 52. Dhossche DM, van der Steen LF, Shettar SM. [Catatonia in autism spectrum disorders: review and case-report]. *Tijdschr Psychiatr*. 2015;57(2):89-93.
- 53. Mahato RS, San Gabriel MC, Longshore CT, Schnur DB. A Case of Treatmentresistant Depression and Body Dysmorphic Disorder: The Role of Electroconvulsive Therapy Revisited. *Innov Clin Neurosci.* 2016;13(7-8):37-40.
- 54. Guo JN, Kothari JS, Leckman JF, Ostroff RB. Successful Treatment of Tourette Syndrome With Electroconvulsive Therapy: A Case Report. *Biological psychiatry*. 2016;79(5):e13-e14.
- 55. Clinebell K, Valpey R, Walker T, Gopalan P, Azzam P. Self-Enucleation and Severe Ocular Injury in the Psychiatric Setting. *Psychosomatics*. 2016;57(1):25-30.
- 56. Katz R, Bukanova E, Ostroff R. Procedural Consolidation During Electroconvulsive Therapy for a Patient With Severe Tourette Syndrome. *The journal of ECT*. 2017;33(1):e7-e8.
- 57. Sajith SG, Liew SF, Tor PC. Response to Electroconvulsive Therapy in Patients With Autism Spectrum Disorder and Intractable Challenging Behaviors Associated With Symptoms of Catatonia. *The journal of ECT*. 2017;33(1):63-67.