

Research Article

Response Rates of rTMS Treatment for Depression at Tranquil TMS - a UK-based Clinic

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Abstract

Background: There is strong evidence supporting efficacy of repetitive Transcranial Magnetic Stimulation (rTMS) in the treatment of depression. There is a paucity of research in the UK in this area.

Aim: The aim of the study was to measure the response rate of rTMS in depression in the UK population.

Methods: Patients attending Tranquil TMS clinic at North West of England from the 1st Feb 2018 to 31st Dec 2019 were included in the study. HDRS and HAM-A applied before and at the end of the treatment to measure the response rate. This was then compared to response rates for similar patient groups from international studies.

Results: This sample showed remission rate of 37.03%. Response rate was 66.66%. Dropout rate was 18.18% which was unrelated to rTMS treatment.

Conclusion: The study once again shows the efficacy of rTMS in the treatment of depression. There is a need of further research in this area in the UK population.

Keywords: Depression; rTMS

Introduction

The treatment of depression has continued to evolve over the years. Despite newer antidepressant medications, the STAR-D trial has clearly shown the limitations of psychopharmacology in the treatment

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of depression [1]. In recent times, neuromodulation has provided a ray of hope for that section of the population whose symptoms fail to respond to traditional measures.

One of these neuromodulation treatments, repetitive Transcranial Magnetic Stimulation (rTMS) has been shown to be an effective treatment for depression. In the UK, rTMS has been approved for the treatment of depression by the NICE in 2015 and the Royal College of Psychiatrists in 2017. In the USA, the FDA has approved this treatment for treatment-resistant depression in 2008 [2]. Health Canada has approved this treatment for the treatment-resistant depression in 2002 [3].

NICE Guidance IPG542 (December 2015) states. The evidence on repetitive transcranial magnetic stimulation for depression shows no major safety concerns. The evidence on its efficacy in the short-term is adequate, although the clinical response is variable. Repetitive transcranial magnetic stimulation for depression may be used with normal arrangements for clinical governance and audit [4]. Studies have shown that repetitive Transcranial Magnetic Stimulation (rTMS) has a definite antidepressant effect on major depression [5]. The evidence suggests that rTMS used solely or in combination with antidepressants for first-episode major depressive disorder may be more effective than antidepressants alone. Thus, the use of rTMS may shorten the treatment odyssey for patients with MDD [6].

Aim

The aim of this study is to evaluate the success rates of the rTMS in the treatment of depression in private North West UK-based clinic.

The general consensus is that 1 in 3 patients achieve complete remission by the end of 1 year and that 1 in 2 patients will show a 50% reduction in their symptoms with the rTMS [7].

We expect patients to halve their scores on two different rating scales -Hamilton Depression Rating Scale (HDRS) and Hamilton Anxiety Rating Scale (HAM-A) by the end of their treatment course (which could last 4-6 weeks) in order to consider it a response to treatment.

For HDRS scores, any score of 7 or under is regarded as remission. The HAM-A scores of 10 or under were regarded as remission for the purpose of this study.

Methods

We analysed data of all the patients who received rTMS treatment in Tranquil TMS from 1st Feb 2018 to 31st Dec 2019. We used 2 rating scales, Hamilton Depression Rating Scale (HDRS) and Hamilton Anxiety Rating Scale (HAM-A).

All patients undertook HDRS and HAM-A assessments at the beginning of the treatment and this was then repeated at the start of every week of their treatment subsequently. For the purposes of this study, we decided to only include the scores from the start of the treatment and following the completion of the treatment.

Results

During the study period, 33 patients commenced rTMS treatment. Out of this number, 6 patients dropped out by the end of the 1st week (18.18%). The reasons for dropouts were varied-2 people thought clinic was geographically too far to commute, 3 patients changed their mind within the first few days regarding the treatment and 1 had a family incident - but none of the dropouts were due to side effect issues.

Our sample included 27 patients all of whom were on at least 1 antidepressant medication and showing no significant response to it. Only 4 patients were drug naive. 27 patients out of 33 (81.81%) completed the treatment see (Table 1) Out of those, 9 patients received intermittent theta bursts protocol (50 Hz) for depression whilst 18 patients received standard protocol for depression (10Hz).

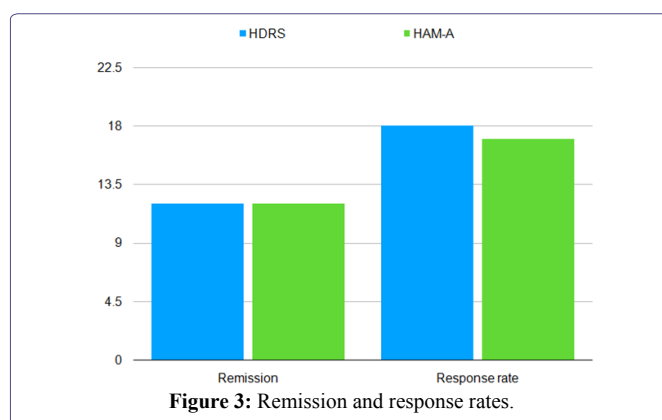
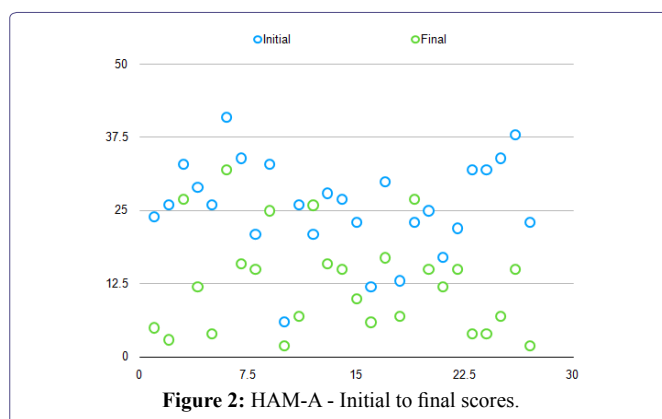
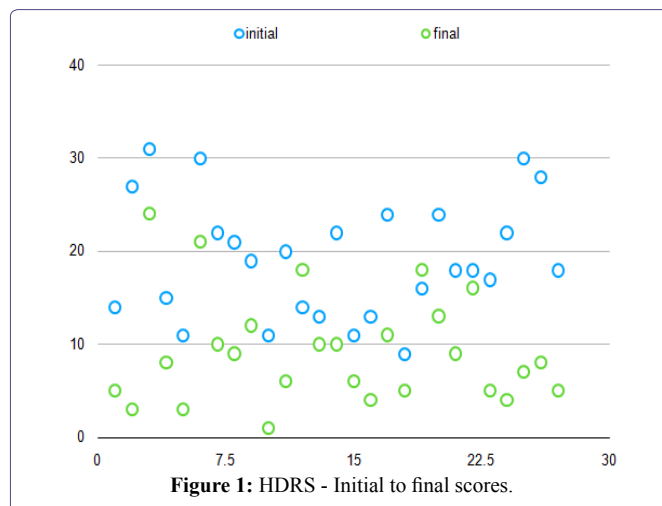
	HDRS initial	HDRS Final	HAM-A Initial	HAM-A final
Patient 1	14	5	24	5
Patient 2	27	3	26	3
Patient 3	31	24	33	27
Patient 4	15	8	29	12
Patient 5	11	3	26	4
Patient 6	30	21	41	32
Patient 7	22	10	34	16
Patient 8	21	9	21	15
Patient 9	19	12	33	25
Patient 10	11	1	6	2
Patient 11	20	6	26	7
Patient 12	14	18	21	26
Patient 13	13	10	28	16
Patient 14	22	10	27	15
Patient 15	11	6	23	10
Patient 16	13	4	12	6
Patient 17	24	11	30	17
Patient 18	9	5	13	7
Patient 19	16	18	23	27
Patient 20	24	13	25	15
Patient 21	18	9	17	12
Patient 22	18	16	22	15
Patient 23	17	5	35	2
Patient 24	22	4	32	4
Patient 25	30	7	34	7
Patient 26	28	8	38	15
Patient 27	18	5	23	2

Table 1: Below highlights the change in HDRS scores from initial to final rating scores.

Patients received 20-40 treatments of high frequency rTMS on the Left Dorsolateral Pre-frontal cortex and a minimum of 10 treatments of low frequency rTMS (1 Hz) on the right dorsolateral prefrontal cortex. Out of the 27, 2 patients (7.4%) are currently receiving maintenance treatment every few weeks

HDRS scores showed 12 out of the 27 patients achieved full remission at the end of the treatment (37.03%). Out of 27 patients 18 had achieved more than a 50% drop in HDRS score (66.66%).

On HAM-A scale see (Figures 1-3) below 12 out of the 27 patients achieved full remission with scores of 10 or under (37.03%). 17 out of 27 patients had achieved more than 50% drop in HAM-A score (62.96%).



Discussion

The study showed that our remission rate for rTMS was high (37.03%). This is comparable to the study done by Connolly et al, 2012 which showed the HDRS remission rate of 35.3% [8].

Our response rate (66.66%) was higher than the response rate of 41.2% quoted in the study by Connolly et al, 2012 [8]. Our rates were also higher than the meta-analysis done by Berlim et al in 2014. Berlim et al carried-out analysis of the data from 29 RCTs covering 1371 patients. They found that 29.3% of patients responded to treatment and 18.6% of patients could achieve remission with high frequency rTMS [9].

We did not have any patients discontinuing the treatment because of the side effects of rTMS. Though our dropout rates were higher than other studies, they were not related to rTMS treatment as such. Once again, this study shows that rTMS is an effective treatment for depression including treatment resistant depression. HAM-A scores highlights the success of anxiety protocol in helping patients with co-existing anxiety symptoms. There is a dearth in the literature regarding rTMS experience in the UK. We recommend further studies with larger sample size mainly targeting the UK's population.

Declaration of Interest

Nikhila Deshpande and Babu Nayar are the directors of Tranquil TMS.

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