



Repetitive Transcranial Magnetic Stimulation (rTMS) Authorization Request Form

Securely email form to: outpatientteam@carelon.com

Please attach your intake assessment for TMS that documents the items below for: diagnosis (and associated symptoms), past trials of TMS, psychotherapy, psychopharmacology, and psychometric measurement.

| | | |
|--|----------------|---------|
| <input type="checkbox"/> In Network | Out of Network | |
| Member Name: | DOB: | Gender: |
| Health Plan: | Policy #: | |
| Date and Time of Request: | | |
| Treating Clinician/Facility: | | |
| If the treating clinician is not making this request, has the treating clinician been notified? <input type="checkbox"/> Yes <input type="checkbox"/> No | | |
| Phone #: | NPI/TIN: | |
| Servicing Clinician/Facility: | | |
| Phone #: | NPI/TIN: | |

1. Diagnosis code and description:

2. Does the Member have a history of TMS attempts in the past?

- ☐ Yes
☐ No

If yes, was there a positive outcome?

- ☐ Yes
☐ No

3. Has the Member had an adequate trial of evidence-based psychotherapy, without significant improvement within the past 5 years?

- ☐ Yes
☐ No

Type of psychotherapy:

Dates of evidence-based psychotherapy trial:

If the Member has not had an adequate trial of evidence-based psychotherapy, what is the reason?

4. Please fill in the Member's psychotropic medications taken within the past five years:

| Medication Name | Dose | Dates of Use (Start and End Dates) | Response Atypical Agents |
|-----------------|------|------------------------------------|--|
| | | | <input type="checkbox"/> Improved <input type="checkbox"/> Inadequate Response <input type="checkbox"/> Adverse Response <input type="checkbox"/> Intolerability <input type="checkbox"/> Non-adherence <input type="checkbox"/> Other _____ |
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(continued)

| Medication Name | Dose | Dates of Use (Start and End Dates) | Response Atypical Agents |
|--|------|------------------------------------|--|
| | | | <input type="checkbox"/> Improved <input type="checkbox"/> Inadequate Response <input type="checkbox"/> Adverse Response <input type="checkbox"/> Intolerability <input type="checkbox"/> Non-adherence <input type="checkbox"/> Other _____ |
| Please list any Augmenting Agents used: _____ | | | |
| If no medications were used, are they contraindicated? | | | |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | | | |
| 5. Were any of these meds used during this depressive episode? | | | |
| <input type="checkbox"/> Yes, list medications: _____ <input type="checkbox"/> No | | | |
| If yes, was improvement inadequate at adequate dose and duration? | | | |
| <input type="checkbox"/> Yes, list dose and duration: _____ <input type="checkbox"/> No | | | |
| If yes, was the medication discontinued due to side effects? | | | |
| <input type="checkbox"/> Yes, list side effects: _____ <input type="checkbox"/> No | | | |
| 6. Please check all that apply: | | | |
| <input type="checkbox"/> Vagus Nerve Stimulator leads in the carotid sheath <input type="checkbox"/> Other implanted stimulators controlled by or that use electrical or magnetic signals <input type="checkbox"/> Conductive or ferromagnetic or other magnetic-sensitive metals implanted or embedded in head or neck within 11.81 inches (30 cm) of TMS coil placement other than dental fillings <input type="checkbox"/> Acute or chronic psychotic disorder <input type="checkbox"/> Seizure disorder or history of seizure disorder <input type="checkbox"/> Substance abuse at time of treatments <input type="checkbox"/> Severe dementia <input type="checkbox"/> Non-adherence with previous depression treatments <input type="checkbox"/> None of the above | | | |
| 7. Will the first treatment session include determining correct magnetic pulse strength and placement of the magnetic coil? | | | |
| <input type="checkbox"/> Yes <input type="checkbox"/> No | | | |
| 8. What is the Member's most recent score on a validated self-report depression rating scale? | | | |
| Rating scale used: | | | |
| Score: | | | |
| Date completed: | | | |