Table 1s Study schedule/procedures

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Protocol Activity** | **Screening** | **Pre-treatment** | **Treatment** | **Post-treatment** | **Follow-up** |
| **Study week** | **-4 - 0** | **1-2** | **3-10** | **11** | **12** |
| **Clinical visit** | **Visit 1 (week 0, baseline)** |  | **Visit 2 (week 10)** |  |  |
| **Phone visit** | **Week 0** | **Week 1** | **weekly** |  | **Week 12** |
| **Informed Consent** | X |  |  |  |  |
| **Medical History & Demographics** | X |  |  |  |  |
| **Physical Exam** | X |  |  |  |  |
| **Inclusion/exclusion criteria** | X |  |  |  |  |
| **Vital signs** | X |  | X X |  |  |
| **Blood collection** | X |  |  |  |  |
| **BPI** | X | X | X |  |  |
| **PROMIS questionnaires** | X |  | X |  |  |
| **QST session** | X |  | X |  |  |
| **AE recording** |  | X | X X | X | X |

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Supplemental Figure 1s: Patients’ perceived therapeutic effect of Gralise versus side effects of Gralise. Patients were categorized as having marked therapeutic relief, mild relief or unchanged/worse pain. Patients were also categorized as having no side effects, not significant and significant side effects.

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Supplemental Figure 2s: Patients’ psychosocial variables, including amount and restfulness of sleep, degree of depression and anxiety, pain catastrophizing, and pain interference with daily functioning were compared between week 1 and week 10 of Gralise therapy.