**SUPPLEMENTAL TABLE 1.** Adverse Events (frequency) during maintenance trial testing lisdexamfetamine and placebo treatments.

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| --- | --- | --- |
|  | **Lisdexamfetamine**  *n*=32  n (%) | **Placebo**  *n*=29  n (%) |
| All-Cause Mortality | 0 (0.0%) | 0 (0.0%) |
| Serious Adverse Events 0 (0.0%) 0 (0.0%)  Adverse Events Leading to Discontinuation\* 2 (6.3%) 1 (3.4%)  Adverse Events (not including Serious)\*\* | | |
| Decreased Appetite | 10 (31.3%) | 4 (13.8%) |
| Dry Mouth | 6 (18.8%) | 1 (3.4%) |
| Insomnia | 6 (18.8%) | 4 (13.8%) |
| Increased Energy | 5 (15.6%) | 3 (10.3%) |
| Constipation | 3 (9.4%) | 1 (3.4%) |
| Jittery | 2 (6.3%) | 0 (0.0%) |
| Increased Heart Rate | 2 (6.3%) | 1 (3.4%) |

Note: Adverse (side effect) events assessed systematically at month one of maintenance randomized double-blind placebo-controlled trial testing lisdexamfetamine versus placebo. These adverse event data are included in clinicaltrials.gov record (NCT03926052).

\* Adverse events leading to discontinuation (medical withdrawal) were reported for 2 cases in lisdexamfetamine (headache, dry eyes) and for 1 case in placebo (increased blood pressure).

\*\* Adverse events reported by >5% of participants in lisdexamfetamine or placebo treatment condition.