**Supplemental Digital Content 11: Ocular and nonocular adverse events up to month 12 suspected to be related to the study drug treatment (Safety set\*)**

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| **Preferred term,  n (%)** | **Ranibizumab 0.5 mg  (n = 119)** | **Sham with ranibizumab  0.5 mg (n = 52)** | **Total (N = 178)** |
| **Ocular AEs, total** | **2 (1.7)** | **1 (1.9)** | **3 (1.7)** |
| Retinal cyst | 1 (0.8) | 0 | 1 (0.6) |
| Vitreous floaters | 1 (0.8) | 0 | 1 (0.6) |
| RPE tear | 0 | 1 (1.9) | 1 (0.6) |
| IOP increased | 1 (0.8) | 0 | 1 (0.6) |
| **Nonocular AEs, total** | **1 (0.8)** | **0** | **1 (0.6)** |
| Arrhythmia | 1 (0.8) | 0 | 1 (0.6) |
| Blood pressure increased | 1 (0.8) | 0 | 1 (0.6) |
| \*Consisted of all adult patients who received at least one application of study treatment and had at least one post-baseline safety assessment.  No AEs were reported for patients in the sham without ranibizumab group.  Preferred terms are sorted in descending order of frequency of the ranibizumab 0.5 mg group. A patient with multiple occurrences of a preferred term was counted only once in the preferred term row.  AE, adverse event; IOP, intraocular pressure; RPE, retinal pigment epithelium | | | |