***Instructions: Review all new or unresolved adverse events (currently on the Adverse Event Log in the DREAM clinical database) with the patient. If the Serious Event Type is greater than 1, complete a Serious Adverse Event Initial Reporting Form for the first report or a Serious Adverse Event Follow-up Form for new additional information. If this is an ocular adverse event involving one eye, please check the R or L boxes; if this is an ocular adverse event involving both eyes check both the R and L boxes.***

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|  | **Adverse Event Coding (NIH)** | **DREAM AE Coding** |
| **AE # (Record from database)** | **MedDRA Code** | **MedDRA Short Name** | **MedDRA Grade** | **Was Event Serious?****No Yes** | **Serious Event Type** | **Tx for AE****No Yes** | **Out-****come** | **DREAM****Tx** | **Ocular AE Eye(s)****R L** | **Start Date****(MMDDYYYY)** | **Stop Date****(MMDDYYYY)** |
| **aeseqg** | **aecode** | **aeterm** | **aesev** | **aeser** | **aeserious** | **aetx** | **aeout** | **aedrtx** | **reye****leye** | **aestdtc** | **aeendtc** |
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| **Completed by: qclname Certification #: qcconcert Date Completed: qccompdtc** |