ATRIAL FIBRILLATION (AFIB)/A-FLUTTER

All Classes (Updated 05/29/2024)

DISEASE/CONDITION	EVALUATION DATA	DISPOSITION
A. Previously reported to FAA and the airman has a letter from the FAA that monitoring is not required.	The airman should bring his/her letter(s) from the FAA (for this condition) for the AME to review. If the AME's history and exam do not reveal any evidence or concern of recurrence:	ISSUE Summarize this history in Block 60.
B. Previously warned;	Submit the following to the FAA for	
Now with New event	review:	DEFER
or Findings:	 Non-Valvular Atrial Fibrillation (AFib)/A-Flutter Status Summary AND Detailed clinical progress notes from the treating physician(s) describing the new event or finding. PLUS: Current ≥ 24-hour cardiac monitor. 	Submit the information to the FAA for a possible Special Issuance. Follow-up Special Issuance – Will be per the Airman's authorization letter
C. Non-Valvular	Submit the following to the FAA for	
AFib/A-Flutter History of at any time OR current: Single or multiple episodes Paroxysmal Persistent Permanent/chronic Untreated or treated AFib treated with ablation (3-month recovery period) or cardioversion (1-month recovery period)	review: Non-Valvular Atrial Fibrillation (AFib)/A-Flutter Status Summary AND Detailed clinical progress notes from any provider seen for this condition (PCP, cardiology, urgent care) which identify symptoms, testing performed, diagnosis, and treatment. Hospital records (if applicable), including: Admission (H&P) Discharge summary Current ≥ 24-hour cardiac monitor Initial etiology work-up as follows: TSH; Sleep Study that meets current AASM or CMS Guidelines for a Type I	Submit the information to the FAA for a possible Special Issuance. Follow-up Special Issuance — Will be per the Airman's authorization letter See Non-Valvular Atrial Fibrillation (AFib)/A-Flutter Status Summary

DISEASE/CONDITION	EVALUATION DATA	DISPOSITION
	(Type III or Type IV NOT allowed); ○ Cardiac echocardiogram; and ○ Exercise stress test □ If taking warfarin, submit info listed on Pharmaceutical Anticoagulants – Emboli Mitigation.	
D. Treated with	After a 6-month recovery period,	
left atrial appendage	submit the following to the FAA for	DEFER
ex: Watchman	review: Cardiologist evaluation that describes why the procedure/device was indicated, treatment regimen throughout the process, any procedure complications, whether device is working properly, and the current status of AFib;	Submit the information to the FAA for a possible Special Issuance. Follow-up Special Issuance – Will be per the Airman's authorization letter
	 □ Current CHA2DS2-VASc score; □ Current ≥ 24-hour cardiac monitor □ Initial AFib etiology work up (TSH, sleep study that meets current AASM or CMS Guidelines for a Type I or Type II sleep study [Type III or Type IV not allowed], cardiac echocardiogram, exercise stress test), if not previously submitted; □ Procedure report; 	
	 □ TEE report from time of implantation, if performed (images not required in most cases); and □ TEE report from ≥ 45 days post procedure to evaluate for peridevice leaks (Recommended images at 0, 45, 90, and 135 degrees with 2-4 heartbeats to show appendage and occlusion device or in accordance with industry standards). 	