VirtuOx Holter Transition to MCT or CEM Criteria Form

less than 5 hours of interpretable data is recorded in the first 24-hour period, VirtuOx will default to the next 24-hour period for determination.

Arrhythmia	Rate and Duration	Change Instructions
Bradycardia (all rhythms other than AF)	< 40 BPM if Symptomatic **OR** < 40 BPM **&** > 30 Seconds if Asymptomatic	
Narrow Complex (Non-Sinus) Tachycardias	> 10 Beats if > 150 BPM ***OR*** > 60 Seconds at Any Rate	
New Onset Atrial Fibrillation / Flutter	> 15 seconds **OR** any episode if symptomatic	
Pause	≥ 4 Seconds if asymptomatic or ≥ 2 Seconds if symptomatic	
Ventricular Tachycardia	≥ 10 beats **OR** any episode if symptomatic	
Accelerated Junctional Rhythm	HR > 40 bpm	
PVC Burden	> 10% of total beats	
2nd Degree AVB, Type II	All Episodes	
3rd degree / Complete AVB	All Episodes	
Ventricular Fibrillation / Toursades de Pointes TdP / Agonal	All Episodes	
Asystole > 6 Seconds	All Episodes	

This criteria applies to (Check all that apply and specify details if necessary)

☐ Physician ☐ Practice ☐ Location(s) ☐ Patient

Additional instructions:

This criteria authorizes VirtuOx's cardiology department to enact a proactive care study if a patient's Holter monitor is non-diagnostic based on the arrhythmia criteria outlined on this form. If the study is unremarkable per this arrhythmia criteria then the study will be optimized to a Mobile Cardiac Telemetry test for 7-days unless a different length has been specified in written physician orders. A Comprehensive report will then be provided to the ordering physician.

Physician name: _____

Physician signature: _____ Date: _____