Signal-averaged electrocardiography (SAECG) is a technique involving computerized analysis of small segments of a standard EKG to detect abnormalities, termed ventricular late potentials (VLP), that would be otherwise obscured by “background” skeletal muscle activity.

Signal-averaged electrocardiography (SAECG) is considered not medically necessary for all indications, including but not limited to the following:

A. Assessment of efficacy of antiarrhythmia drug therapy
B. Assessment of success after surgery for arrhythmia
C. Assessment of success of pharmacological, mechanical, or surgical interventions to restore coronary artery blood flow
D. Cardiomyopathy
E. Detection of acute rejection of heart transplants
F. Risk stratification for ventricular arrhythmia following acute myocardial infarction
G. Risk stratification of patients with Brugada syndrome
H. Syncope

NOTE: A summary of the supporting rationale for the policy criteria is at the end of the policy.

CROSS REFERENCES
None

BACKGROUND

VLPs reflect aberrant, asynchronous electrical impulses arising from viable isolated cardiac muscle bordering an infarcted area, and are thought to be responsible for ventricular tachyarrhythmias. Therefore, VLPs, as measured by SAECG, have been investigated as a risk factor for arrhythmic events in patients with a variety of cardiac conditions, including cardiomyopathy and prior history of myocardial infarction (MI).

Patients considered at high risk of ventricular arrhythmias, and thus sudden death, may be treated with drugs to suppress the emergence of arrhythmias or implantable cardiac defibrillators (ICD) to promptly detect and terminate tachyarrhythmias when they occur. Because sudden cardiac death, whether from arrhythmias or pump failure, is one of the most common causes of death after a previous MI, there is intense interest in risk stratification to target therapy. The focus of this policy is on primary prevention in patients who have not experienced a life-threatening arrhythmia and who may benefit from treatment.

VLP is just one of many risk factors that have been investigated. Others include left ventricular ejection fraction, arrhythmias detected on Holter monitor or electrophysiologic studies, heart rate variability, and baroreceptor sensitivity. T-wave alternans is another technique for risk stratification; it measures beat-to-beat variability, while SAECG measures beat-averaged conduction.

EVIDENCE SUMMARY

In a clinical area such as cardiac rhythm abnormalities where multiple tools to predict risk already exist, use of signal-averaged electrocardiography (SAECG) must demonstrate that any improvement in predictive accuracy results in meaningful changes in therapy and leads to improved outcomes. In many cases, comparative trials are needed to demonstrate the impact of testing on net health outcomes.

CLINICAL VALIDITY

SAECG has been studied as a risk stratification tool for potentially fatal arrhythmias in patients with a previous myocardial infarction (MI). Studies have failed to demonstrate SAECG’s ability to accurately identify patients at risk for sudden cardiac death. Positive predictive values (i.e., the ability of the test to identify patients who will experience ventricular arrhythmias) were low (8-44%) and varied between studies, depending on the population studied. Negative predictive values (i.e., the ability of the test to identify patients who will not experience ventricular arrhythmias) were high (88-97%), but it has not been demonstrated that this information is helpful in the overall clinical management of the patient. However, a key statistic underlying the negative predictive value is the underlying prevalence of the outcome. Although sudden cardiac death is the most common cause of death in the one-year period after...
infarction, it is relatively uncommon (2.5–11.3%) and declining as a result of increasing use of thrombolytic therapy, aspirin, and beta-blockers. Thus, given the relative low incidence of arrhythmias, the high negative predictive value is not surprising.

CLINICAL UTILITY

The ultimate validation of any diagnostic test is to determine how it is used in the management of patients and whether the management results in improved health outcomes. SAECG has not been successfully used as a patient selection criterion in the clinical randomized trials investigating both drug and device antiarrhythmic therapy in the post MI patient. Also, no study definitively reported a decrease in fatal arrhythmias as a direct result of using SAECG for risk stratification and subsequent treatment decisions. Published studies have failed to demonstrate SAECG’s ability to impact clinical management.

SAECG, used as a risk stratification tool, either showed no improvement in survival or proved to be only a weak predictor of sudden cardiac death.[4-11]

The CABG-Patch trial recruited patients scheduled for a CABG who had an ejection fraction of less than 36% and abnormalities on the SAECG.[12] AECG was not used alone as a risk stratification tool in this study. Patients were randomized to a defibrillator group or a control group and all received CABG. There was no evidence of improved survival among those in the defibrillator group. However, it cannot be determined whether the failure of this trial was due to the selection criteria or the treatments being compared. No conclusions can be drawn about the utility of SAECG in determining the patient’s course of clinical management.

Results of SAECG were found to be a weak predictor of sudden cardiac death in a nonrandomized consecutive series of 700 patients with a history of acute MI.[4] These results are unreliable due to the nonrandomized study design.

A small controlled clinical trial observed a correlation of various markers that identified patients with Brugada syndrome who were at risk for life-threatening arrhythmias.[7] Late potentials identified on SAECG appeared to be the most useful for identifying patients potentially at risk for ventricular fibrillation and sudden cardiac death.

An accompanying editorial identified the study limitations and methodological details that required further clarification.[8] Each patient did not receive all of the risk stratification tools being compared. The authors stated that, even though this is a rare disease, the study population was too small to establish statistical significance. It was unknown if patients were taken off of sodium channel blockers or if SAECG was measured only on unpaced complexes. Although results of the study suggested a role for SAECG as a risk stratifier, there was no clear evidence that the test would predict which patients would become symptomatic and which would not.

SAECG was evaluated in a study using an algorithm for risk stratification to determine appropriateness for prophylactic ICD implantation.[13] The algorithm also included left ventricular ejection fraction, programmed ventricular stimulation, and family history of sudden cardiac death. While results were promising, only 69 patients received SAECG and larger, randomized studies are needed to confirm the clinical utility of SAECG in risk-stratifying algorithms.

PRACTICE GUIDELINE SUMMARY
A 2009 updated consensus document by the American College of Cardiology/American Heart Association (ACC/AHA) recommended against routine use of SAECG in adults with heart failure because it “has not been shown to provide incremental value in assessing overall prognosis” in these patients.[14,15] This was a class III recommendation, defined as a procedure that should not be performed as it is not helpful and may be harmful; no additional studies are needed.

## SUMMARY

The current research shows that signal-average electrocardiography (SAECG) has not been used successfully to determine and stratify patients into clinically relevant categories of risk. In addition, there is no research that states the use of SAECG can be used to guide patient management and no clinical guidelines based on research recommend SAECG for any indication. Therefore, SAECG is considered not medically necessary for all indications.

## REFERENCES


### CODES

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