Novel Electrophysiology Signal Recording (n) **System Enables Specific Visualization** of the Purkinje Network and **Other High-Frequency Signals**



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igh-frequency signals (e.g., Purkinje potentials) are of interest in arrhythmia syn-L dromes such as fascicular tachycardia and ventricular fibrillation. However, restricted dynamic range and sampling rate in modern electrophysiology recording systems make it challenging to identify high-frequency, low-amplitude signals, especially those temporally situated near low-frequency, highamplitude signals or large sharp peaks. Although increasing gain accentuates their appearance, clipping and artifact may occur.

It is undetermined whether: 1) increased dynamic range and sampling rate; and 2) use of additional filtering and processing (PURE-EP, BioSig Technologies, Minneapolis, Minnesota), will improve recording of Purkinje potentials and other high-frequency signals. As such, we performed unipolar and bipolar cardiac conduction system mapping using various catheters while simultaneously comparing PURE-EP (0.05 to 1 kHz, 24-bit A/D conversion, 2000 samples/s sampling rate, low gain with no saturation in the ± 250 mV range) with a standard recording system (0.05 to 500 Hz, 12-bit A/D conversion, 977 samples per second sampling rate). The additional high-pass filtering and processing in PURE-EP enables the detection

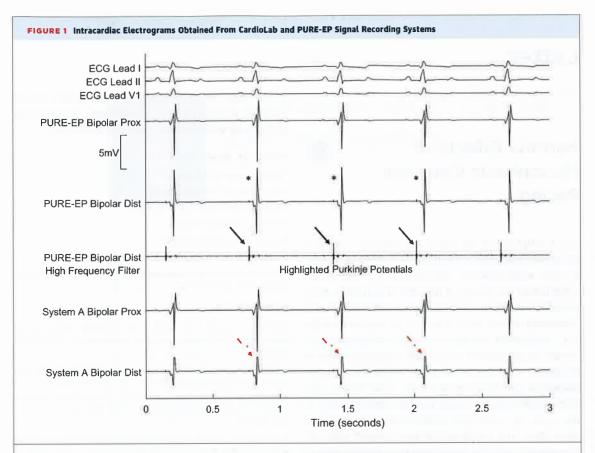
and visualization of high-frequency signals in the presence of larger waveforms.

PURE-EP consistently demonstrated superior Purkinje potentials and other high-frequency signal visualization compared with the standard recorder in all sites of the cardiac conduction system. The PURE-EP system has the capability to display the same channel with different processing options to highlight specific features while still displaying the original electrogram signal. For example, the new system only demonstrated high-frequency signals which correlated with Purkinje potentials (Figure 1). Traditional EP recording systems have fixed dynamic range and sampling rate and, therefore, lack the adjustability demonstrated by the PURE-EP system. Such features may improve mapping/ablation outcomes in arrhythmia syndromes dependent on the Purkinje

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Compared with CardioLab, PURE-EP demonstrated high-frequency signals without artifact and clipping (*). In addition, the PURE-EP automated filtering algorithm illustrates high-frequency signals that correlated with Purkinje potentials (black arrows). Although CardioLab demonstrates the Purkinje signals, clipping of the ventricular electrogram occurs (red arrows).

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