



Natus Receives FDA Clearance for the NeuroLink 256 Channel Amplifier

- New Device Extends Range of Neurodiagnostic Solutions

SAN CARLOS, Calif. (June 10, 2010) – **Natus Medical Incorporated** (Nasdaq:BABY) today announced the regulatory clearance and immediate North American market availability of its NeuroLink IP 256-channel amplifier. The clearance was granted to Natus by the FDA through the 510(k) application process.

Introduction of a high-density 256-channel amplifier is an important step in the non-invasive localization of partial epileptic seizures. Studies show a relationship between the number of electrodes used in an EEG exam and the accuracy of localizing brain abnormalities, which is important in the diagnosis and treatment of seizure disorders.

With the addition of the 256-channel NeuroLink IP amplifier to its existing offering of 64- and 128-channel networkable amplifiers, Natus continues its tradition of bringing innovative solutions into the epilepsy market. With this offering, Natus becomes the only manufacturer with a brain monitoring product line that ranges from a single-channel aEEG device to the 256-channel NeuroLink IP.

About Natus Medical

Natus is a leading provider of healthcare products used for the screening, detection, treatment, monitoring and tracking of common medical ailments in newborn care, hearing impairment, neurological dysfunction, epilepsy, sleep disorders, and balance and mobility disorders. Product offerings include computerized neurodiagnostic systems for audiology, neurology, polysomnography, and neonatology, as well as newborn care products such as hearing screening systems, phototherapy devices for the treatment of newborn jaundice, head-cooling products for the treatment of brain injury in newborns, and software systems for managing and tracking disorders and diseases for public health laboratories.

Additional information about Natus Medical can be found at www.natus.com.

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