

SciBase receives US patent for Nevisense electrode design

May 2nd SciBase received patent approval for a further US patent.

On May 2nd 2017 SciBase received confirmation that US patent number 9,636,035 has been approved. The patent protects the unique shape of the microneedles that enable detection of malignant melanoma in the skin using electrical impedance as demonstrated with high precision in clinical trials. In total SciBase now has 3 patent families approved in the US and an additional family in process.

“Our electrode design is one of the key reasons for the success of our method. It is positive that we further strengthen our patent portfolio in the US, especially as we are in the final stages of our PMA-application.”, says Simon Grant CEO SciBase.

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This information is information that SciBase Holding AB is obliged to make public pursuant to the EU Market Abuse Regulation. The information was submitted for publication, through the agency of the contact person set out above, at 08.00 CET on May 4, 2017.

About Skin Cancer

Skin cancer is one of the most common cancers in the world, accounting for nearly half of all cancers. It has been estimated that nearly half of all Americans who live to the age of 65 will develop skin cancer at least once. Malignant melanoma is the most fatal form of skin cancer causing the majority (75%) of deaths related to skin cancer. Worldwide, doctors diagnose about 230,000 new cases of melanoma yearly.

About SciBase and Nevisense

SciBase AB is a Swedish medical technology company, headquartered in Stockholm that has developed a unique point-of-care device for the accurate detection of malignant melanoma. Its product, Nevisense, helps doctors to detect malignant melanoma, the most dangerous type of skin cancer. SciBase was founded by Stig Ollmar, Associate Professor at The Karolinska Institute in Stockholm, Sweden. Nevisense is based on substantial research and has achieved excellent results in the largest clinical study ever conducted on the detection of malignant melanoma. Nevisense is CE marked in Europe, has TGA approval in Australia, and is awaiting FDA clearance in the United States. Nevisense is based on a method called Electrical Impedance Spectroscopy (EIS), which uses the varying electrical properties of human tissue to categorize cellular structures and thereby detect malignancies. SciBase is listed on Nasdaq First North (“SCIB”). Avanza is the certified advisor. Further information is available on www.scibase.com.