US FDA 510(k) clearance for the Biodenta Tapered Implant System

March 2013 - Good news from the Biodenta Bone Level Tapered Implant System registration!

In March 2013, Biodenta received 510(k) clearance from the US Food and Drug Administration ("FDA") to market the Biodenta Dental Implant System "Bone Level Tapered".

The 510(k) number is: K123415

This clearance allows Biodenta to register, import and distribute the Biodenta Bone Level Tapered products into the US market.

David Eiler, Quality Manager of Biodenta Group, commented: "FDA clearance for the Bone Level Tapered Implant Line is once more a very important step in the product development. I shall like to thank everybody, who was involved in this project, especially for the support regarding clinical publications, document compiling, and updating the surgical and prosthetic guidelines".

Read more about the Biodenta Bone Level Tapered Implant System