



V9

Free Sales Certificate

The Danish Medicines Agency hereby certifies that the medical devices specified in the attached list are manufactured by:

Sanispaces ApS
Hammerholmen 39B
2650 Hvidovre
Denmark

Medical devices which are CE marked in conformity with Directive 93/42/EEC meet the essential requirements for safety and performance. They may therefore be manufactured and marketed in Denmark and exported without any approval from the Danish Medicines Agency.



Valid from: 18 November 2020
Valid Until: 18 November 2022

Lasse Nielsen



List of products registered with The Danish Medicine Agency:

- Sanispace 60H
- Sanispace 125
- Sanispace 125T
- Sanispace 250



LÆGEMIDDELSTYRELSEN
DANISH MEDICINES AGENCY