Ajuvia™/ Perspectis, Inc.
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EUROPEAN DECLARATION OF CONFORMITY

We, **Perspectis Inc.**
1 First Canadian Place, Ste. 350,
Toronto, ON, Canada, M5X 1C1,

declare under our sole responsibility

that the product: **BACK VITALIZER**

of risk class: 1 (according to rule 1 set out in annex VIII of Regulation (EU) 2017/745)

CE-marked with CE

to which this declaration relates is in full conformity with the following

- COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices
  - REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC

Authorized Representative: Loox Company e.K.
Sedanstr. 26
30161 Hanover, Germany

The Technical Construction File

is maintained at: 1 First Canadian Place, Ste. 350, Toronto, ON, Canada, M5X 1C1.

This declaration of conformity is valid until December 31st, 2026.

Date of issue: January 1st, 2021

Place of issue: Toronto, Canada

Patrick Lee
President, Perspectis Inc.