

DECLARATION OF CONFORMITY UE

Name of Manufacturer: **Blackfin Spa, Via Nogarola, 17 32027 Taibon Agordino (BL)** ,
in its capacity as the manufacturer hereby declares, under its sole responsibility, that the optical
frame:

MODELS			
941	AERO A-N	1015	VECTOR TWO
942	AERO A-M	1069	AERO LOOP B1-P1
943	AERO VECTOR	1070	AERO LOOP B1-P2
943	VECTOR ONE	1071	AERO LOOP B2-P1
947	FISHBOURNE	1072	AERO LOOP B2-P2
948	HAYLE	1073	AERO LOOP B3-P1
957	BEMBRIDGE	1074	AERO LOOP B3-P2
958	ALTON	1075	AERO LOOP B4-P1
959	ST. HELEN	1076	AERO LOOP B4-P2

in all its skus (colors and sizes)

is a class I medical device (Rule 1 Chapter III of Annex VIII of (EU) Regulation 2017/745), which is
intended for use in connection with, in the manufacturing of eyeglasses, ophthalmic lenses for
correcting vision problems.

We also declare that it is in conformity with the provisions of (EU) Regulation 2017/745 on
Medical Devices and with the standard EN ISO 12870:2018

BASIC UDI-DI CODE: 803277450CQ02100201B3

Taibon Agordino 31/07/2024

Blackfin, created and hand made in Italy by
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