Accreditation **SCESp 0046**
Notified Body **1247**
Accredited Certification Authority
in accordance with ISO/IEC 17065:2012

NSBIV AG
Certification body
SIBE Schweiz

EU type-examination certificate

Translation of original EU type-examination certificate in German

No. 5004/1

Identification of PPE Category III, g)	Lanyard according EN 354:2010
Brand	AirWork & Heliseilerei GmbH (A&H)
Type	A, B, C or D Configuration of types according application instruction: AWA_A&H-PSA_sPCDS Approved components according: 260107_A&H-PSA_ReZert-EU-BMB_5004-1_Artikelliste_A
Additional information about the PPE	Lanyard according EN 354:2010 (black / anthracite or black / yellow) with or without metal connections with lenght from 10 cm up to 200 cm. Designed for transport or securing of a person during helicopter operations. Dynamic forces from falls are not permitted.
Fully or partially applied harmonised standards, technical specifications	EN 354:2010
Address of manufacturer	AirWork & Heliseilerei GmbH (A&H) Bahnhofweg 1 CH-6405 Immensee
Valid until	13 January 2031

The verified PPE type against falling from a height complies with the essential health and safety requirements of Regulation (EU) 2016/425, Annex II. This certificate may only be used in combination with the procedure of internal production control with supervised product checks at random intervals (Module C2) in accordance with Annex VII (Annex V, 6.2, item j, k). This certificate is only being validity together with any enclosures mentioned on front page as well as the legal provisions listed on the reverse side.

Date of issue
14 January 2026

P. Müller
Safety Engineer

NSBIV AG
Certification Body *SIBE Schweiz*
Brünigstrasse 18
CH-6005 Lucerne

R. Walker
Head of Certification Body

Legal provisions

(1) This EU type-examination certificate is valid only for personal protective equipment (PPE) against falling from a height (hereafter named for PPE type) with the brand and type specification mentioned on front page. Each PPE against falling from height shall be marked according Annex VII, fig. 6, to certify compliance with Regulation (EU) No 2016/425 on PPE.

(2) The EU type-examination certificate may have one or more annexes attached.

(3) The notified body shall keep itself appraised of any changes in the generally acknowledged state of the art which indicate that the approved type may no longer comply with the applicable essential health and safety requirements and shall determine whether such changes require further investigation. If so, the notified body shall inform the manufacturer accordingly.

(4) The manufacturer shall inform the notified body that holds the technical documentation relating to the EU type-examination certificate of all modifications to the approved type and of all modifications of the technical documentation that may affect the conformity of the PPE with the applicable essential health and safety requirements or the conditions for validity of this certificate. Such modifications shall require additional approval in the form of an addition to the original EU type-examination certificate.

(5) The manufacturer shall ensure that the PPE continues to fulfil the applicable essential health and safety requirements in light of the state of the art. Moreover, he shall take all measures necessary so that the manufacturing process and its monitoring ensure conformity of the manufactured PPE with the type described in the EU type-examination certificate and with the applicable requirements of this Regulation.

(6) The manufacturer shall ask the notified body to review the EU type-examination certificate either:
(a) in the case of a modification to the approved type referred to in point (4);
(b) in the case of a change in the state of the art referred to in point (5);
(c) at the latest, before the date of expiry of the certificate.
In order to allow the notified body to fulfil its tasks, the manufacturer shall submit his application at the earliest 12 months and at the latest 6 months prior to the expiry date of the EU type-examination certificate.

(7) The notified body shall examine the PPE type and, where necessary in the light of the changes made, carry out the relevant tests to ensure that the approved type continues to fulfil the applicable essential health and safety requirements. If the notified body is satisfied that the approved type continues to fulfil the applicable health and safety requirements, it shall renew the EU type-examination certificate. The notified body shall ensure that the review procedure is finalised before the expiry date of the EU type-examination certificate.

(8) Where the conditions referred to in points (a) and (b) of point (6) are not met, a simplified review procedure shall apply. If the notified body finds that a change in the state of the art referred to in point (5) has occurred, the procedure set out in point (7) shall apply.

(9) If, following the review, the notified body concludes that the EU type-examination certificate is no longer valid, the body shall withdraw it and the manufacturer shall cease the placing on the market of the PPE concerned.

(10) The Commission, the Member States and the other notified bodies may, on request, obtain a copy of the EU type-examination certificates and/or additions thereto. On a reasoned request, the Commission and the Member States may obtain a copy of the technical documentation and the results of the examinations carried out by the notified body.

(11) The notified body shall keep a copy of the EU type-examination certificate, its annexes and additions, as well as the technical file including the documentation submitted by the manufacturer, for a period of five years after the expiry of the validity of that certificate. The manufacturer shall keep a copy of the EU type-examination certificate, its annexes and additions, together with the technical documentation at the disposal of the national authorities, for 10 years after the PPE has been placed on the market.

(12) Before placing PPE on the market, the manufacturer shall lodge an application for supervised product checks at random intervals with a single notified body of his choice. The requirements for the supervised product check at random intervals are set out in Annex VII of Regulation (EU) No 2016/425 on PPE (Module C2).

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