

**EU DECLARATION OF CONFORMITY
REGULATION (EU) 2016/425**

We: Syntech International, Inc.
17171 Daimler Street
Irvine, CA 92614 USA
Phone Number: 1 (800) 443-3842
www.syntech-intl.com

Declare under sole responsibility, that our products were evaluated by **BSI Group The Netherlands B.V.**, Notified Body 2797 and deemed them to be compliant with Annex V (Module B) of the Personal Protective Equipment Regulation (EU) 2016/425 and the applicable Essential Health & Safety Requirements.

Declare under sole responsibility that our products conform to the type Based on Quality Assurance of the production process (Module D) under surveillance **BSI Group The Netherlands B.V.**, Notified Body 2797

Compliance with the applicable Essential Health & Safety Requirements has been demonstrated thru examination in accordance with the harmonized standard,

EN 12941: 1998 +A2: 2008

Product description: Respiratory protective device – Powered filtering device incorporating a helmet or hood.

Models: CAPR System(s)

Certificate Reference # CE 813956
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Category: III (three)

Classification: TH2

Filters: The following filters/filter combinations are permitted:

Filter	Type / Class
2167-10 XP	TH2 P R S
2166-10 OV	TH2 P R S

DLC Cuffs

Part reference	Part number	Description
2365-02SM	01031316	DLC Lens-Cuff SM
2365-02ML	01031291	DLC Lens-Cuff ML
2081-03	03531001	Helmet-Ch Universal
2071-08	03531021	Helmet Liner
2061-08	01031284	Filter Cover Cap

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DLC Shroud

Part reference	Part number	Description
2260-05 SM	01031435	DLC Shroud SM
2260-05 ML	01031434	DLC Shroud ML
2081-3	03531001	Helmet-Ch Universal
2071-08	03531021	Helmet Liner
2061-08	01031284	Filter Cover Cap

Battery options:

Part reference	Part number	Description
2500-36TSC	01532104	Lithium Ion Battery 14.4V, 2.25Ah
2500-37TSC	01532161	Lithium Ion Battery (Extended Capacity) 14.4V, 3.35Ah
2000-76	-	Lithium Ion Battery Belt
2600-01	01432089	Li Battery Charger
2600-02	01432202	Li Battery Charger
2590-05	-	Power Cord

The CAPR can be used with either battery or charger.

We hereby appoint mdi Europa GmbH, Langenhagener Str. 71 30855 Hannover-Langenhagen, Germany to act as EU Authorized Representative as explicitly defined in article 9 of the **REGULATION (EU) 2016/425 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 9 MARCH 2016**

Authorised Signatory:

Eric Fiskewold

Eric Fiskewold (May 13, 2025 11:17 PDT)

Eric Fiskewold /
Quality Assurance Manager

May 13, 2025

13 MAY 2025

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CE-Marking Experts for Medical Devices

Langenhagen, 5 June 2025

SYNTECHINTL

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F22004 REV F

F22004 REV F_Declaration of Conformity, BSI

Final Audit Report

2025-05-13

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