

## Combination Agreement

The medical equipment/devices manufactured by ALBER in the table below can be combined with the equipment/devices manufactured by Alu Rehab AS in the table below, provided that the combination is carried out in a professional manner and in accordance with the restrictions of both manufacturers, as indicated in the user manuals and installation instructions, provided by the manufacturers.

Wheelchair Model Device Name	Alber Device								Remarks
	e-ifx E35/E36	e-motion M25 & Duo Drive	viamobil V14	viamobil V25	via GO V24	e-pilot P15	smoov O10	Scalamobil S35	
Netti V	✓	✓	–	✓	–	–	–	–	Including Base, All-round
Netti II/III	✓	✓	–	✓	✓	–	–	✓	Including EL, HD
Netti 4U	✓	✓	–	✓	✓	–	–	✓	Including CE, CE Plus, CED, Base
Netti Dynamic (Adapt Pro)	✓	✓	–	–	–	–	–	–	
Sense	✓	✓	–	–	–	–	–	–	

Combination of the **ALBER** power assist with **Alu Rehab AS** wheelchairs must be executed according to the ALBER mounting instructions by authorized and trained staff only. Alber restrictions (e.g. seat width, maximum user weight) have to be observed (check compatibility online with Alber [Dealer Information Pool / DIP](#) before starting the process of ordering and combining).

Observe the product limitations and instructions for use given in the user manuals of all equipment mentioned in this combination agreement. Machining, bending, welding, or bracing of safety relevant components is not permitted. Exceptions are permitted in accordance with the manufacturer's installation instructions.

ALBER shall be responsible for the verification activities, including the relevant documentation relating to the combination. The relevant documentation shall be made available to Alu Rehab AS in a timely manner upon request.

Both parties shall communicate planned changes to their device as far as the combination may be affected.

All external communication, in particular communication to the competent authorities regarding the combination, shall first be communicated between the parties. In general, communication to the authorities shall be made by the manufacturer of the device which has caused an adverse event.

This agreement is valid from date of signature and is subject to three months' notice for termination. If regulatory requirements should change or if constructional changes are made to the products, each of the parties shall evaluate whether this affects the validity of this agreement and inform the other party without any delay of such circumstances.

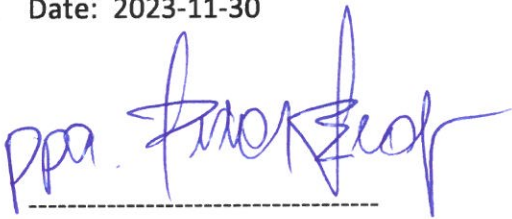
Support for requests from authorities and requests for investigation of adverse events shall be provided by both parties within the timeframe specified by the authority or manufacturer.

This Combination Agreement was issued in duplicate and signed duly by authorised representatives.

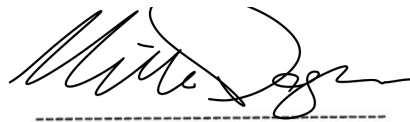
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Date: 2023-11-30

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