

TANGO

Integrated pediatric and adult spine boards



Tango spine boards are devices designed to lift and immobilize patients with suspected spinal injuries.

Specific features

The integration of adult and pediatric spinal boards, offers unmatched versatility

Pediatric spinal board with differentiated occipital areas for the best adaptation to the anatomical characteristics of pediatric patients

The pediatric spinal board can be easily extracted from the adult one, ensuring fast rescue operations and minimizing storage space

The head immobilizer can easily be placed thanks to strap bands

Technical data

Spinal board dimensions	1830 x 445 x 55 mm
Number of handles	14
Pediatric spinal board weight	3 kg
Adult spinal board weight	8 kg
Weight of complete spinal board	11 kg
Tango load capacity	150 kg
Baby Go load capacity	30 kg
Pediatric spinal board dimensions	1190 x 320 x 45 mm
Pediatric spinal board number of handles	10
Materials	Polyethylene

Accessories

ST02102B	Spine Pack – Carry bag
ST02106A	MARK UP – Permanent personalization system
ST02105A	Fix Board – 10 G Wall support for spinal board

Class I MD compliant with UE Reg. 2017/745

TANGO ADULT AND PAEDIATRIC SPINE BOARD



ST02140C

CND Classification V0804

Registration number 101670

Rev.0 (14/06/2021)

UNCHECKED COPY – further revisions will be available on <http://support.spencer.it>

Spencer Italia S.r.l. Sala Baganza (PR) Italia Tel. +39.0521.541111 Fax +39.0521.541222

www.spencer.it

EU DECLARATION OF CONFORMITY/ DICHIARAZIONE DI CONFORMITA' UE
Regulation/Regolamento UE 2017/745

The declaration is released under the sole responsibility of the manufacturer
La dichiarazione è rilasciata sotto la responsabilità esclusiva del fabbricante

Manufacturer/*Fabbricante*: **Spencer Italia s.r.l.**
Via Provinciale, 12 – 43038 Sala Baganza (PR) – Italy

Unique registration number/
Numero di registrazione unico: IT-MF-000027507

Medical Device/*Dispositivo Medico*: TANGO ADULT AND PAEDIATRIC SPINE BOARD/
TANGO TAVOLA SPINALE.PEDIATR.E ADULTO INTEGR.

Code/*Codice*: ST02140C

BASIC UDI-DI /*UDI-DI di base*: 805771123TAVOLESPINALITT

Lot/ *Lotto* SN/ *Matricola*: Not available before the production/
Non disponibile prima della produzione

Quantity/*Quantità*: 1

Risk class /*Classe di rischio*: I
(Annex VIII/*Allegato VII*)

Conformity assessment procedure/
Procedura valutazione conformità: Not present/Non presente

Rule/*Regola*: 1

Spencer Italia S.r.l. declares under its sole responsibility that the above mentioned medical device is in compliance with the requirements of the Regulation 2017/745 and with the applicable regulations and common specifications.

Spencer Italia S.r.l. dichiara sotto la sua sola responsabilità che il Dispositivo Medico sopra menzionato, è conforme ai requisiti del Regolamento 2017/745, alle norme e alle specifiche comuni applicabili.

The list of applicable standards is reported in the Technical File.
La lista delle norme applicabili è riportata nel relativo Fascicolo Tecnico.

Sala Baganza (PR) - IT, 11/01/2023

First name and surname /*Nome e cognome*: _____

Role/*Ruolo*: _____ Signature/*Firma*: _____

Person in whose name and on whose behalf this declaration of conformity has been signed/
Persona a nome e per conto della quale è stata firmata la presente Dichiarazione UE :

Antonio Ciardella
(Legal Representative /*Legale Rappresentante*)



CERTIFICAT

CERTIFICADO

СЕРТИФИКАТ

認證證書

CERTIFICATE

ZERTIFIKAT



Italia

CERTIFICATO

Nr. 50 100 6189 Rev.006

SI ATTESTA CHE / THIS IS TO CERTIFY THAT

IL SISTEMA DI GESTIONE PER LA QUALITÀ DI
THE QUALITY MANAGEMENT SYSTEM OF

Spencer Italia s.r.l.

SEDE LEGALE E OPERATIVA:
REGISTERED OFFICE AND OPERATIONAL SITE:

**VIA PROVINCIALE 12
IT - 43038 SALA BAGANZA (PR)**

SEDI OPERATIVE: VEDI ALLEGATO 1 / OPERATIONAL SITES: SEE ANNEX 1

È CONFORME AI REQUISITI DELLA NORMA
HAS BEEN FOUND TO COMPLY WITH THE REQUIREMENTS OF

UNI EN ISO 9001:2015

QUESTO CERTIFICATO È VALIDO PER IL SEGUENTE CAMPO DI APPLICAZIONE
THIS CERTIFICATE IS VALID FOR THE FOLLOWING SCOPE OF APPLICATION

Progettazione, sviluppo e produzione di dispositivi per l'emergenza e funerario. Commercializzazione di prodotti per il settore medicale, emergenza, soccorso e funerario a marchio proprio e non (IAF 19, 14, 29)

Design, development and production of emergency and mortuary equipment. Distribution of medical, emergency rescue and mortuary equipment with own and other brands (IAF 19, 14, 29)



SGQ N° 049A

Membro degli Accordi di Mutuo Riconoscimento
EA, IAF e ILAC
Signatory of EA, IAF and ILAC Mutual
Recognition Agreements

Per l'Organismo di Certificazione
For the Certification Body
TÜV Italia S.r.l.

Validità / Validity

Dal / From: **2021-08-05**

Al / To: **2024-08-04**

Data emissione / Issuing Date

Andrea Coscia
Direttore Divisione Business Assurance
Business Assurance Division Manager

2021-08-05

PRIMA CERTIFICAZIONE / FIRST CERTIFICATION: 2006-10-03

DATA DI SCADENZA DELL'ULTIMO CICLO DI CERTIFICAZIONE: 2021-08-04
EXPIRATION DATE OF THE LAST CERTIFICATION CYCLE: 2021-08-04

"LA VALIDITÀ DEL PRESENTE CERTIFICATO È SUBORDINATA A SORVEGLIANZA PERIODICA A 12 MESI E AL RIESAME COMPLETO DEL SISTEMA DI GESTIONE AZIENDALE CON PERIODICITÀ TRIENNALE"
"THE VALIDITY OF THE PRESENT CERTIFICATE DEPENDS ON THE ANNUAL SURVEILLANCE EVERY 12 MONTHS AND ON THE COMPLETE REVIEW OF COMPANY'S MANAGEMENT SYSTEM AFTER THREE-YEARS"



Italia

ALLEGATO 1 AL CERTIFICATO NR 50 100 6189 Rev.006**ANNEX 1 TO CERTIFICATE NO 50 100 6189 Rev.006**

pagina 1 di 1 / page 1 of 1

IL CERTIFICATO NR 50 100 6189 Rev.006 COPRE ANCHE LE SEGUENTI SEDI OPERATIVE:
THE CERTIFICATE N 50 100 6189 Rev.006 COVERS ALSO THE FOLLOWING OFFICES:**Spencer Italia s.r.l.****VIA PROVINCIALE 12
IT - 43038 SALA BAGANZA (PR)**

Progettazione, sviluppo e produzione di dispositivi per l'emergenza e funerario. Commercializzazione di prodotti per il settore medicale, emergenza, soccorso e funerario a marchio proprio e non

*Design, development and production of emergency and mortuary equipment.
Distribution of medical, emergency rescue and mortuary equipment with own and other brands***VIA PETITOT 4
IT - 43038 SALA BAGANZA (PR)**

Immagazzinamento di prodotti per il settore medicale, emergenza e soccorso a marchio proprio e non

*Warehousing of medical, emergency and rescue equipment with own and other brands***VIA LEGA DEI CARRETTIERI 3
IT - 43038 SALA BAGANZA (PR)**

Immagazzinamento di prodotti per il settore medicale, emergenza e soccorso a marchio proprio e non

*Warehousing and Distribution of medical, emergency and rescue equipment with own and other brands***VIA PROVINCIALE 38
IT - 43038 SALA BAGANZA (PR)**

Immagazzinamento di prodotti per il settore medicale, emergenza e soccorso a marchio proprio e non

Warehousing of medical, emergency and rescue equipment with own and other brands

SGQ N° 049A

Membro degli Accordi di Mutuo Riconoscimento
EA, IAF e ILAC
Signatory of EA, IAF and ILAC Mutual
Recognition AgreementsPer l'Organismo di Certificazione
For the Certification Body
TÜV Italia S.r.l.

Validità / Validity

Dal / From: **2021-08-05**Al / To: **2024-08-04**
Andrea Coscia
Direttore Divisione Business Assurance
Business Assurance Division Manager

Data emissione / Issuing Date

2021-08-05**PRIMA CERTIFICAZIONE / FIRST CERTIFICATION: 2006-10-03**

DATA DI SCADENZA DELL'ULTIMO CICLO DI CERTIFICAZIONE: 2021-08-04

EXPIRATION DATE OF THE LAST CERTIFICATION CYCLE: 2021-08-04

"LA VALIDITÀ DEL PRESENTE CERTIFICATO È SUBORDINATA A SORVEGLIANZA PERIODICA A 12 MESI E AL RIESAME COMPLETO DEL SISTEMA DI GESTIONE AZIENDALE CON PERIODICITÀ TRIENNALE"
"THE VALIDITY OF THE PRESENT CERTIFICATE DEPENDS ON THE ANNUAL SURVEILLANCE EVERY 12 MONTHS AND ON THE COMPLETE REVIEW OF COMPANY'S MANAGEMENT SYSTEM AFTER THREE-YEARS"



Certificate

No. Q5 033230 0033 Rev. 00

Holder of Certificate: **Spencer Italia s.r.l.**
Via Provinciale 12
43038 Sala Baganza (PR)
ITALY

Certification Mark:



Scope of Certificate: **Design and development, production of emergency equipment; Distribution of medical, emergency and rescue equipment with own and other brands**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:Q5 033230 0033 Rev. 00](http://www.tuvsud.com/ps-cert?q=cert:Q5_033230_0033_Rev.00)

Report No.: ITA 1683049

Valid from: 2021-11-09
Valid until: 2024-08-04

Date, 2021-11-09

Christoph Dicks
Head of Certification/Notified Body

Certificate

No. Q5 033230 0033 Rev. 00

Applied Standard(s):

EN ISO 13485:2016
Medical devices - Quality management systems -
Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

Facility(ies):

Spencer Italia s.r.l.
Via Provinciale 38, 43038 Sala Baganza (PR), ITALY

Warehousing of medical, emergency and rescue equipment with
own and other brands.

Spencer Italia s.r.l.
Via Provinciale 12, 43038 Sala Baganza (PR), ITALY

Design and development,, production of emergency equipment;
Distribution of medical, emergency and rescue equipment with
own and other brands

Spencer Italia s.r.l.
Via Petitot 4, 43038 Sala Baganza (PR), ITALY

Warehousing of medical, emergency and rescue equipment with
own and other brands.

Spencer Italia srl
Via Lega dei Carrettieri 3, 43038 Sala Baganza (PR), ITALY

Warehousing and Distribution of medical, emergency and rescue
equipment with own and other brands.

\