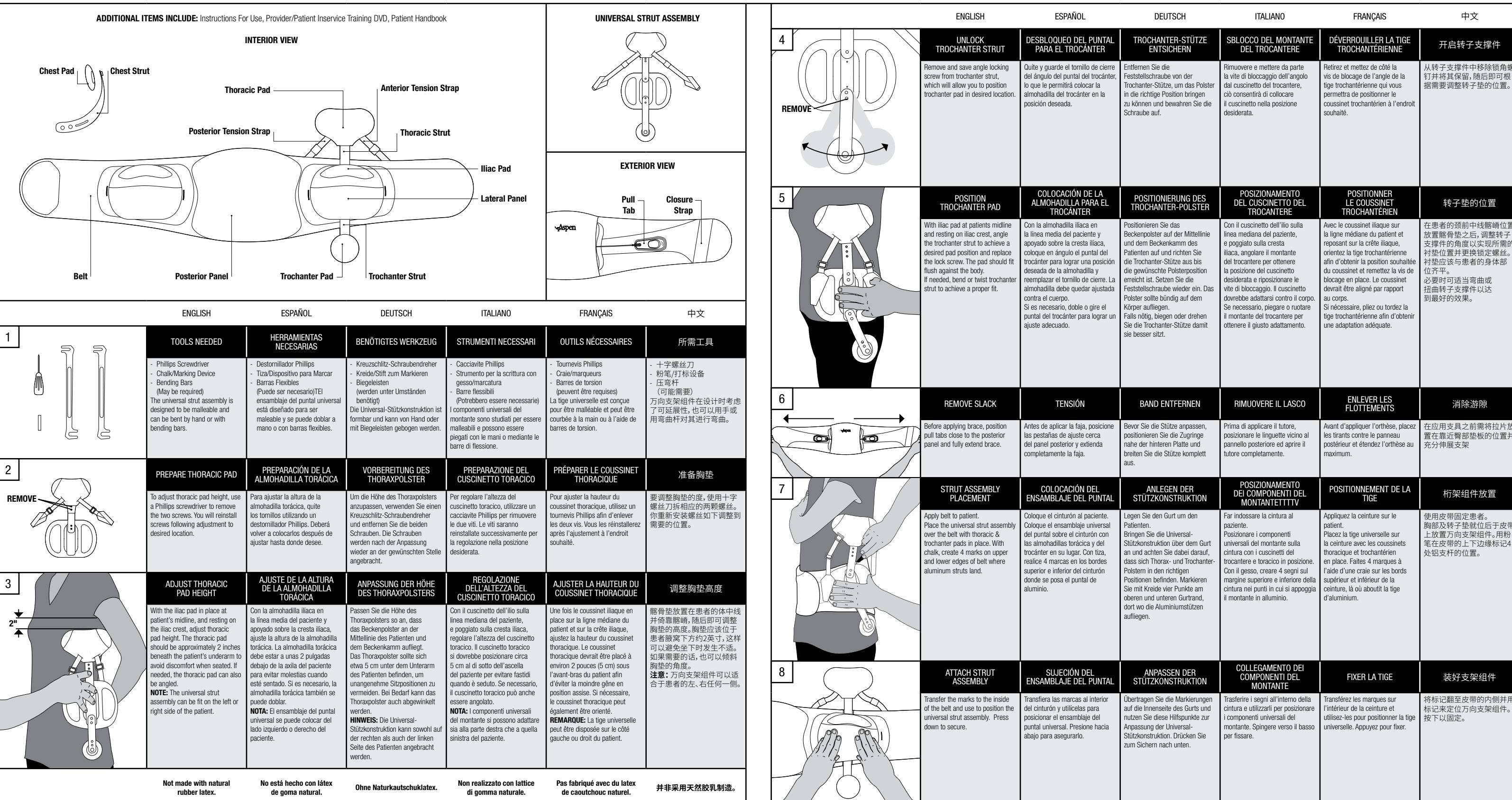


Peak Scoliosis Bracing System™

INTENDED FOR ADULT PATIENTS
DUAL CONFIGURATION KIT AVAILABLE
FOR BI-LATERAL CONFIGURATION

ASPEN MEDICAL PRODUCTS
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	ENGLISH	ESPAÑOL	DEUTSCH	ITALIANO	FRANÇAIS	中文		ENGLISH	ESPAÑOL	DEUTSCH	ITALIANO	FRANÇAIS	中文
9		ATTACH TENSION STRAPS	SUJECCIÓN DE LAS CORREAS DE TENSIÓN	ANBRINGEN DER SPANNGURTE	COLLEGAMENTO DELLE CINGHIE DI TENSIONAMENTO	ATTACHER LES SANGLES DE TENSION	调整胸垫高度		TIGHTEN AND ADJUST	APRIETE Y AJUSTE	FESTZIEHEN UND ANPASSEN	STRINGERE E REGOLARE	SERRER ET AJUSTER
10		PLACE LATERAL PANEL	COLOCACIÓN DEL PANEL LATERAL	ANBRINGEN DER SEITENPLATTE	POSIZIONAMENTO DEL PANNELLO LATERALE	PLACER LE PANNEAU LATÉRAL	调整胸垫高度		CONFIRM COMFORT	CORROBORACIÓN DE VLA COMODIDAD	SICHERSTELLUNG DES KOMFORTS	VERIFICA DEL COMFORT	VÉRIFIER LE CONFORT
11		APPLY THE BRACE	COLOCACIÓN DE LA FAJA	ANBRINGEN DER STÜTZE	APPLICAZIONE DEL TUTORIO	APPLIQUER L'ORTHÈSE	调整胸垫高度		REMOVING BRACE	RETIRO DE LA FAJA	ENTFERNEN DER STÜTZE ENTFERNEN	RIMOZIONE DEL TUTORE	ENLEVER L'ORTHESE
12		COMPRESS BELT	COMPRESIÓN DEL CINTURÓN	FESTZURREN DES GURTS	COMPRESSE DELL'AFLATURA	COMPRESSER LA CEINTURE	调整胸垫高度		IDENTIFY CORRECT SIDE	IDENTIFICACIÓN DEL LADO CORRECTO	IDENTIFIZIERUNG DER RICHTIGEN SEITE	IDENTIFICACION DEL LATO CORRETTO	IDENTIFIER LE CÔTÉ APPROPRIÉ
13		A	COMPRESS BELT	COMPRESIÓN DEL CINTURÓN	FESTZURREN DES GURTS	COMPRESSE DELL'AFLATURA	COMPRESSER LA CEINTURE		INSTALL CHEST PAD	IDENTIFICACIÓN DEL LADO CORRECTO	IDENTIFIZIERUNG DER RICHTIGEN SEITE	POSIZIONAMENTO DEL CUSCINETTO PETTORALE	INSTALLER LE COUSSINET PECTORAL
14		B	NOTE: Instruct patient to apply brace in the same position each time.						POSITION CHEST PAD	COLOCACIÓN DE LA ALMOHADILLA PARA EL PECHO	POSITIONIERUNG DES BRUSTPOLSTERS	POSIZIONAMENTO DEL CUSCINETTO PETTORALE	POSITIONNER LE COUSSINET PECTORAL
SIZING	SMALL	MEDIUM	LARGE	EXTRA LARGE									调整胸垫位置
CIRCUMFERENCE	24 – 30 in 61 – 76 cm	29 – 35 in 74 – 89 cm	34 – 40 in 86 – 102 cm	39 – 45 in 99 – 114 cm									

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(b) The product is altered;

(c) Any parts not supplied by Aspen Medical Products are inserted into the product; or

(d) The product is not used in accordance with the Aspen Medical Products Instructions for Use.

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