Instructions for Use

REF Premium Head & Chin Straps 800-0369

INTENDED USE
Intended use is to hold patient's head during operative procedure. The intended users of this device are medical professionals within hospitals and surgery centers.

INSTRUCTIONS
Become familiar with features of patient positioning device before use with patient. Always practice use on a nurse, physician or appropriate volunteer prior to using clinically.

1. *Place patient on surgical table/shoulder chair in supine position*
2. *Once patient is intubated, hold patient's head/neck and raise shoulder chair up so patient is in Fowler (sitting) position*
3. *Loosen ball joint - adjust head positioner around patient's head making sure patient's ears are uncovered - tighten ball joint*
4. *Hold head positioner flaps alongside of patient's head, placing Forehead Strap across forehead as shown. Attach securely and avoid covering patient's eyebrows.*
5. *Center Chin Strap over chin, secure strap to one side of head positioner. Making sure head is properly positioned, secure strap's other end to opposite side of head positioner.*

NOTE: Head should be securely positioned. If not, make straps a little tighter.

COMPONENT OVERVIEW
Premium Head and Chin Straps holds a patient's head securely during operative procedure.

GENERAL SPECIFICATIONS
Device Dimensions (maximum)

- *Length: 21"+/- 0.5" (53 cm +/- 1 cm) (Head)*
- *Length: 22"+/- 0.5" (56 cm +/- 1 cm) (Chin)*
- *Width: 2"+/- 0.5 (5 cm +/- 1 cm) (Both Head and Chin)*
- *Device Weight: .066 lbs. +/- .02 lbs. (.03 kg +/- .009 kg)*
- *Attaches to head section of SchureMed Shoulder Chairs*
- *Single-person installation*

GENERAL INFORMATION

- *Product not made with Natural Rubber Latex*

- *If damaged in shipping, call Customer Service at (888) 724-8763 or (781) 982-7000 to obtain Return Material Authorization number. For product warranty issues, contact Customer Service.*
- *CE marked medical device according to MDR (EU) 2017/745*
- *Store device between -4°F to +86°F (-20°C to 30°C)*

SINGLE PATIENT USE—Please dispose after each use

USER NOTICE
Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

eIFU Language Versions
To download and print the Instructions for Use, please go to www.schuremed.com

Symbol Glossary

Symbol	Title
	Manufacturer
	Date of manufacture
EC REP	Authorized Representative in the European Community
CH REP	Authorized Representative in the Swiss
	Importer
SN	Serial Number
	Warning
MD	Medical Device
UDI	Unique Device Identifier
CE	CE Marking
	Single Patient Use

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