

Brun Medical Co., Ltd.

GRA-407002,Rev.02

## Connecting Tubing [INSTRUCTION FOR USE]

### 1. INTENDED PURPOSE

The product is used for connecting pipelines and devices which provides pathway of the liquid-injection or/and invasive blood pressure monitoring in interventional operation by professional physician.

### 2. INTENDED USER

Intended users are the competent physicians who have the training in interventional treatment.

### 3. INTENDED PATIENT POPULATION

Patients needing interventional treatment.

### 4. CONTRAINDICATION

No absolute contraindications when used correctly by medical personnel.

### 5. HANDING AND STORAGE

Stored in the relative humidity of not more than 80%, non-corrosive gases, a common, dry, well-ventilated and clean environment.

### 6. STRUCTURE AND SPECIFICATIONS

The device is consist of the male and female luer connectors , tubing.

Specifications:

Specifications	material	OD(mm)	ID (mm)	Length(cm)	Pressure ( psi )
BR-CT1.80	PVC	1.80	0.80	10、 20、 30、 60、 90、 120、 150	500
BR-CT2.00	PVC	2.00	1.00	10、 20、 30、 60、 90、 120、 150	500
BR-CT2.70	PVC	2.70	1.30	10、 20、 30、 60、 90、 120、 150	500
BR-CT3.00	PVC	3.00	1.50	10、 20、 30、 60、 90、 120、 150	500
BR-CT3.20	PVC	3.20	1.50	10、 20、 30、 60、 90、 120、 150	500
BR-CT3.20A	PVC	3.20	1.50	20、 60、 90、 120、 150、 180、 200	1000
BR-CT3.65	PVC	3.65	2.00	10、 20、 30、 60、 90、 120、 150	500
BR-CT3.65B	PU	3.65	2.00	20、 60、 90、 120、 150、 180、 200	1200
BR-CT4.00	PVC	4.00	3.00	10、 20、 30、 60、 90、 120、 150	500
BR-CT4.00-150S	PVC	4.00	3.00	150	500
BR-CT4.00-150D	PVC	4.00	3.00	150	500

## 7. PRODUCT PERFORMANCE

Flexible tubing and rotating male luer provide easy and fast connection. High pressure transparent material ensures the need for large flow.

## 8. INSTRUCTION FOR USE

1. Remove the packaging, do not use if the package damaged.
2. Attach the connecting tube to luer connector of the related product, and can be used directly after tightening.
3. Single use only, discards after use.

## 9. PRODUCT LABEL AND PACKAGING (See the label or packaging)

1. Product name, model and specification.
2. Production date, batch lot.
3. Sterilization expiry date.

## 10. CAUTION

1. Procedure must be performed by trained medical personnel.
2. Sterilization is valid for three years, please confirm date of sterilization before use, do not use if expiry.
3. The device is designed and intended for SINGLE USE ONLY.
4. Do not re-sterilize and/or reuse, otherwise patients could get the risk of infection.
5. Sterilized by ethylene oxide. Do not use if the package damaged.



Caution



Use-by date



Keep dry



Keep away from sunlight



Catalogue number



Date of manufacture



Manufacturer



Do not re-use



Sterilized Using  
ethylene oxide



Do not use if package is damaged  
and consult instruction for use



Batch code



Unique device identifier



Authorized representative  
in the European Community/  
European Union



Consult instructions for  
use or consult electronic  
instructions for use



Do not re-sterilize



Medical device



Single sterile barrier system



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