



THE TOP IN QUALITY
for all areas of healthcare

TOP INFUSION PUMP MJ-210



Automatic
IV set Attaching Function

Easy
Operation

Specifications

Name	TOP Infusion Pump				
Model	MJ-210				
Power supply	AC adapter : Input 100 ~ 240V±10% 50/60Hz / Output DC 5V 2A Internal battery : Rechargeable DC 3.7V 2500mAh Li-ion Battery Battery life : Approx. 4 hours				
Input current	AC Power supply : 0.9A				
Applicable IV sets	Recommendation	TOP Solution Infusion Set U series			
	Available	General Infusion Set			
Flow rate range	U series mode	0.1 ~ 99.9mL/h (0.1mL/h increments) 100 ~ 1200mL/h (1mL/h increments)			
	General mode	15drops/mL Infusion set : 1 ~ 400mL/h (1mL/h increments) 20drops/mL Infusion set : 1 ~ 360mL/h (1mL/h increments) 60drops/mL Infusion set : 1 ~ 150mL/h (1mL/h increments)			
Volume limit range	0.1 ~ 9999.9 mL (0.1mL increments)				
Total volume range	0.0 ~ 9999.9 mL (0.1mL increments)				
Infusion accuracy	U series mode	±5% (ISO60601-2-24, Clinical situations, water or saline)			
	General mode	±10% (ISO60601-2-24, Clinical situations, water or saline)			
Occlusion detection pressure		Downstream occlusion			Upstream occlusion
		U series mode		General mode	---
	Oc.Lv.3 :	High	120±30kPa	High	0 ~ -100kPa
	Oc.Lv.2 :	Medium	90±30kPa	Medium	
	Oc.Lv.1 :	Low	60±30kPa	Low	
	Oc.Lv.0 :	Lowest	30±20kPa	Lowest	
Air-in-line detection sensitivity	Ar.Lv.0 :	High	(Approx. 50µL)		
	Ar.Lv.1 :	Medium	(Approx. 0.1mL/15minutes)		
	Ar.Lv.2 :	Low	(Approx. 1mL/15minutes)		
Alarm / Warning	High priority alarm	Power loss / System error / Battery off / Volume limit / Air-in-line / Occlusion(Downstream/Upstream) / Drop sensor / Free flow / Eject / Empty container / Flow error(Over/Under)			
	Low priority alarm	Pre-volume limit / No action / No battery / low battery / Unspecified flow rate / Set value confirmation / Data reset			
	Information	Maintenance timer / Power supply switch			
Special functions	Automatic Infusion Set Attaching function / Automatic clamp confirmation function / Anti-bolus function / KVO function / Standby function / Auto power off function / Repeat alarm / Buzzer setting / Power supply switch notice / Key-lock function / Maintenance timer / History function / Alarm setting confirmation function				
Environmental conditions	Ambient temperature : 10 ~ 40°C Relative humidity : 20 ~ 90% (Noncondensing) Atmospheric pressure : 70 ~ 106kPa				
Service life	6 years (based on self certification using TOP Corporation data)				
Classification	Class II  and internally powered equipment, Type CF  , IPX2 (Drip proof II), Continuous operation				
Dimension	180(W) x 113(H) x 202(L) mm				
Weight	Approx. 1.8kg (Including Clamp and Battery)				
Accessories	AC adapter x1 Internal battery x1 Plug(BF or C-2 type) x1 Operation guide x1				
Standard	ISO60601-2-24				

ASAL
ORIGINAL

PIHAK BERKUASA
PERANTI PERUBATAN



MEDICAL DEVICE
AUTHORITY

PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY
AKTA PERANTI PERUBATAN 2012 (AKTA 737)
MEDICAL DEVICE ACT 2012 (ACT 737)
SIJIL PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CERTIFICATE
Seksyen 5(1) Akta 737
Section 5(1) of Act 737

No. Pendaftaran: **GC6722519-36743**
Registration No.:

Tarikh Sah Pendaftaran:
Registration Validity Date:

28/11/2024 - 27/11/2029

Sijil ini adalah dengan ini diberi kepada: **MEDITOP CORPORATION (MALAYSIA) SDN. BHD.**
This certificate is hereby issued to:

yang beralamat di:
which is located at:

**NO. 3, PERSIARAN USAHAWAN, TAMAN IKS,
SEKSYEN 9, ,
43650 BANDAR BARU BANGI
SELANGOR DARUL EHSAN**

bagi mengesahkan peranti perubatan seperti yang dinyatakan dalam Lampiran 1 adalah berdaftar di bawah Seksyen 5(1) Akta 737.

to confirm that the medical device as detailed out in Attachment 1 is registered under Section 5(1) of Act 737.

Pendaftaran ini diberikan tertakluk kepada peruntukan-peruntukan di bawah Akta 737 dan peraturan-peraturan yang dibuat dibawahnya serta syarat-syarat seperti di Lampiran 2.

This registration is granted subject to the provisions under Act 737 and its subsidiary legislations and the conditions as in Attachment 2.



MURALITHARAN PARAMASUA
KETUA EKSEKUTIF
CHIEF EXECUTIVE
PIHAK BERKUASA PERANTI PERUBATAN
MEDICAL DEVICE AUTHORITY



No. Pendaftaran: **GC6722519-36743**
Registration No.:

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Butir-butir peranti perubatan yang didaftarkan
Particulars of the registered medical device

Nama Peranti Perubatan **INFUSION PUMP**
Medical Device Name

Kelas
Class

CLASS C

Jenama
Brand

TOP

Kelompok
Group

SINGLE

Produk Identifikasi
Product Identifier

MJ-210

Nama dan alamat
pembuat:
*Name and address of
manufacturer*

**TOP CORPORATION
19-10 SENJU NAKAI-CHO, ADACHI-KU, TOKYO, JAPAN,
120-0035
JAPAN**



SYARAT – SYARAT PENDAFTARAN PERANTI PERUBATAN
MEDICAL DEVICE REGISTRATION CONDITIONS

1.0 SYARAT AM
GENERAL CONDITIONS

- 1.1 Syarat-syarat pendaftaran peranti perubatan ini dibuat adalah berdasarkan kepada Seksyen 7 (1), Akta Peranti Perubatan 2012 (Akta 737). Kelulusan ini diberi berdasarkan maklumat-maklumat yang telah diterima.
Medical device registration conditions are prescribed in accordance to Section 7(1) of Medical Device Act (Act 737). Approval is granted based on information received.
- 1.2 Establismen hendaklah mematuhi segala arahan yang dikeluarkan oleh Pihak Berkuasa dari semasa ke semasa.
Establishment must comply with all instructions issued by the Authority from time to time.
- 1.3 Pihak Berkuasa berhak untuk meminda syarat-syarat pendaftaran dari semasa ke semasa.
The Authority reserves the rights to amend the registration conditions from time to time.
- 1.4 Pihak Berkuasa berhak menjalankan pemeriksaan ke atas establismen pada bila-bila masa tanpa dimaklumkan terlebih dahulu.
The Authority reserves the right to conduct inspection at any time without prior notice.
- 1.5 Pihak Berkuasa boleh membatalkan Pendaftaran Peranti Perubatan atau mengambil tindakan undang-undang sekiranya Establismen gagal mematuhi mana-mana syarat Pendaftaran Peranti Perubatan.
The Authority may cancel the Medical Device Registration or take legal action if the Establishment fails to comply with any medical device registration conditions.
- 1.6 Sijil Pendaftaran Peranti Perubatan yang dikeluarkan oleh Pihak Berkuasa tidak boleh dipindah milik.
Medical Device Registration Certificate issued by the Authority shall not be transferable or assignable.
- 1.7 Sijil Pendaftaran Peranti Perubatan hendaklah dikemukakan sekiranya diminta oleh mana-mana pegawai yang diberi kuasa.
Medical Device Registration Certificate must be presented upon request by any authorized officer.
- 1.8 Establismen tidak boleh membenarkan Sijil Pendaftaran Peranti Perubatan disalahgunakan oleh individu/syarikat lain dalam apa-apa cara.
Establishment shall not permit the Medical Device Registration Certificate to be abused in any way by any individual / another party.
- 1.9 Tempoh sahlaku Sijil Pendaftaran Peranti Perubatan adalah lima (5) tahun dari tarikh pendaftaran melainkan jika pendaftaran itu dibatalkan oleh Pihak Berkuasa sebelum habis tempohnya.
The validity of the Medical Device Registration Certificate is five (5) years from the date of registration unless the registration is cancelled by the Authority before its expiry.
- 1.10 Sijil ini tidak mengecualikan mana-mana keperluan perundangan lain yang terpakai untuk sesuatu peranti perubatan (contoh: Peranti Perubatan yang mengandungi racun berjadual tertakluk kepada Akta Racun 1952; peranti perubatan menggunakan sinaran mengion adalah tertakluk kepada Akta Perlesenan Tenaga Atom 1984.)
This certificate does not exempt any other regulatory requirements applicable to the medical device (for examples: Medical Device containing scheduled poison is subjected to the Poisons Act 1952; medical devices using ionizing radiation is subjected to the Atomic Energy Licensing Act 1984) .

1.11 Establismen hendaklah melaporkan insiden melibatkan peranti perubatan yang didaftarkan kepada Pihak Berkuasa seperti tertakluk di bawah Seksyen 40 Akta 737.
Establishment shall report any incidents involving registered medical device to the Authority as prescribed in Section 40 of Act 737.

1.12 Peranti perubatan yang diniatkan bagi kegunaan professional hanya boleh dibekalkan untuk kegunaan professional perubatan sahaja dan tidak boleh diletakkan dipasaran bagi kegunaan orang awam.
Medical device intended for professional use may only be supplied for use by medical professionals only and shall not be placed in the market for general public.

2.0 PINDAAN PENDAFTARAN PERANTI PERUBATAN AMENDMENT OF MEDICAL DEVICE REGISTRATION

2.1 Sebarang pindaan kepada maklumat yang berkaitan peranti perubatan yang berdaftar hendaklah dimaklumkan kepada Pihak Berkuasa secara rasmi mengikut garis panduan yang ditetapkan oleh Pihak Berkuasa. Pihak Berkuasa berhak memberikan kelulusan atau menolak permohonan pindaan tersebut.
Any amendments to the information concerning registered medical device shall be notified to the Authority in accordance to the guidelines set by the Authority. The Authority reserves the right to grant approval or reject the application for such amendments.

3.0 PEMBATALAN SIJIL PENDAFTARAN PERANTI PERUBATAN CANCELLATION OF MEDICAL DEVICE REGISTRATION CERTIFICATE

3.1 Sijil Pendaftaran Peranti Perubatan boleh dibatalkan seperti yang dinyatakan dalam Seksyen 9, Akta 737.
Medical Device Registration certificate may be cancelled as prescribed in Section 9 of Act 737.

3.2 Mana-mana peranti perubatan yang dibatalkan Sijil Pendaftarannya, tidak boleh diimport, dieksport atau diletakkan dalam pasaran.
Any Medical Device which the registration certificate has been cancelled shall not be imported, exported or placed in the market.

3.3 Sijil Pendaftaran Peranti Perubatan boleh dibatalkan jika Wakil Diberi Kuasa ditamatkan lantikan oleh pembuat.
Medical Device Registration Certificate may be cancelled if Authorized Representative appointment is terminated by the manufacturer.

4.0 HAK PIHAK BERKUASA THE AUTHORITY OWNERSHIP

4.1 Sijil Pendaftaran Peranti Perubatan yang dikeluarkan fizikal atau maya adalah **Hak Milik Pihak Berkuasa**.
The Authority retains the ownership of every Medical Device Registration Certificate issued by any means.

4.2 Sekiranya berlaku kehilangan atau kerosakan Sijil Pendaftaran Peranti Perubatan, hendaklah dimaklumkan kepada Pihak Berkuasa dan setiap penggantian sijil akan dikenakan caj perkhidmatan.
Any loss or damage to the Medical Device Registration Certificate shall be notified to the Authority and every replacement of certificate shall be liable with service charge rendered.

**5.0 TUGAS DAN TANGGUNGJAWAB
ROLES AND RESPONSIBILITIES**

- 5.1** Establismen hendaklah mematuhi Akta 737, peraturan-peraturan di bawah Akta dan syarat-syarat Pendaftaran Peranti Perubatan.
Establishment shall comply with Act 737, its subsidiary regulations and registration Condition.