



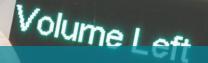


BODYGUARD COLOURVISION 323TM AMBULATORY INFUSION PUMP

The BodyGuard ColourVision 323™ Ambulatory Infusion Pump is a multitherapy pump suitable for parenteral nutrition, homecare environments and hospital infusions with six programmable modes (TPN, continuous, step, intermittent, PCA and drug library).

Main Features:

- Designed to be intuitive to program and use
- Lightweight and compact, suitable for both bedside and ambulatory use
- A delayed start infusion option means no reminder and no additional clinical intervention needed once pump has been set up for general infusion
- The piston pump mechanism ensures unparalleled accuracy and linearity of infusion
- Configurable modes of operation which can be set to a speciality TPN, continuous, step, intermittent, PCA and drug library
- Adjustable air-in-line detection. The cumulative ultrasonic counter can be set according to local protocols
- Variable occlusion pressure settings, configurable to the route of infusion/ patient with on screen pre detection graph of increases in downstream line pressure to help detect potential access device problems at an early stage
- Automatic occlusion alarm cancelling and infusion restart on resolution of inadvertent line kinking episodes



Features and Applications

Safety Features:

- Post Occlusion Bolus Reduction System will safely release pressure from the line, following an occlusion, without administering a post occlusion bolus
- Built-in event log automatically records accurate and reliable data on pump
- Fixed or variable code protected programmes that are configured and locked into the pump memory and can only be accessed by authorised personnel or technicians
- Lockable keypad to prevent inadvertent key presses
- Optional programmable drug library reduces clinician error whilst reducing treatment set-up time. Drugs for specific care units can be enabled to easily navigate your relevant drug list

Modes of Operation:

- TPN
- Continuous
- Step
- Intermittent
- PCA
- Drug library

Intended Use:

The BodyGuard ColourVision 323[™] Ambulatory Infusion Pump is designed for infusion of medications or fluids requiring continuous or intermittent delivery at precisely-controlled infusion rates through clinically acceptable routes of administration, including but not limited to:

- Intravenous
- Subcutaneous
- Percutaneous
- Intra-arterial

Common Applications:

- Parenteral nutrition (PN)
- Enzyme replacement therapy
- Immunology
- Maintenence of general fluids or medication

Speak to our team who have the knowledge and experience to support you in choosing the right medical infusion products to meet your requirements.



The BodyGuard ColourVision 323[™] Ambulatory Infusion Pump has a range of dedicated administration sets. These are optional 1.2 and 0.2 micron filter with optional manual priming valve. All sets are non-DEHP¹ PVC plasticized with TOTM², are BSE-TSE³ free, latex free and EVA⁴ free. Length and priming volume are approximate.

For more information on administration sets or for product codes, please visit our website; cmemedical.co.uk. You can also contact our Customer Support team on 01253 206700 or email customersupport@cmemedical.co.uk

References

1. Bis(2-ethylhexyl) phthalate, 2. Tris (2-Ethylhexyl) Trimellitate, 3. Bovine spongiform encephalopathy, 4. Ethylene-vinyl acetate.



Ongoing Support

Clinical Support Network

We cover the UK and work in partnership with healthcare providers to help ensure successful pump conversions and deliver device user training programmes. All our Clinical Support Specialists have backgrounds in areas such as critical care, pain management, theatre and recovery, community and palliative care.

Technical Support

Our Technical Support team comprises of experienced medtech engineers who ensure quality servicing, maintenance and repair of our products. Please contact our team to discuss the different levels of service support available to best suit your needs.

Quality Assurance and Regulatory Support

Our team of experienced Quality Assurance professionals can resolve customer complaints, conduct first line testing (in conjunction with our Technical Support team), whilst managing relations with local regulatory bodies, supplier and quality audits and regulatory approvals.



Regional Field Sales Support

When you want to make informed choices about which products will meet the needs of your service and patients, our team has the knowledge and experience to talk to you, wherever you are in the UK or Ireland.

Customer Support

Whether you are a clinician, an engineer or a patient using CME Medical products or services, our team of experienced customer support professionals are available to deal with your enquiry.

Technical Specification

Device type	Piston pump
Flow rate	PCA - 0ml to 100 ml/hr in 0.1ml, 100ml to 1200ml/hr in 1ml increments All other programs - 0.1ml to 100ml/hr in 0.1ml, 100ml to 1200ml/hr in 1ml increments
Priming rate	1200ml/hr
Total infused volume	PCA - 0.1ml to 1,000ml Continuous, TPN, Intermittent - 0.1ml to 9999ml 25 Steps - 0.1ml to 9999ml for each step Accumulated volume for all steps - up to 10 litres
Total time setting	100 hours
Pump accuracy	+/- 5%
KVO rate	0ml to 20ml/hr
Air sensor	Ultrasonic, adjustable air bubble size Adjustable air bubble size 0.01ml-2ml for single bubble detection Accumulated air-in-line detection 2ml over 15 minutes
Maximum pressure	Adjustable 100mmHg to 1500mmHg (in 10mmHg increments)
Time to alarm for maximum occlusion pressure	1hr and 45min at a rate of 0.1ml/hr, 30min at a rate of 1ml/hr
Power supply	100-240 VAC, 50/60Hz. 0.3A max



Battery	Type CF Equipment, degree of protection against electrical shock class II equipment, IPX3 protection against ingress of water
Battery operation at 125 ml/hr	15 hours (rechargeable)
Battery charging	Automatic when clicked into the charger that is connected to an AC power source. Six hours needed to charge a fully depleted battery
Alarms	Air in line, down occlusion, pump unattended, end program, low battery, end battery, door open, fatal error, lock mode, lock out mode, missing key
Software connectivity	BodyComm (optional) Remote monitoring system (optional)
Classification	Type CF Equipment (degree of protection against electrical shock), Class II
Moisture protection	IPX3
Housing	PC/ABS (fire retardant)
Weight	390g (with battery)
Electrical safety	Complies with EN60601-1 (Medical Electrical Equipment Safety), IEC60601-2-24 (infusion pumps and controllers), IEC60601-1-4 (Programmable Electrical Medical System)
Standards	Manufactured in accordance to ISO 9001
	CE Marked (in accordance with the Medical Devices Directive 93/42/EEC)
EMC	Designed to be in compliance with EN 60601-1 (safety) and IEC 601-1-2 (EMC)

Technical Specification (Continued)

Aviation environment	Pull tests based on FAR (Federal Aviation Regulations) 29, 27, 25, 23 Tested in accordance to RTCA/DO-160 Guidelines
Environmental specifications	Non operating conditions (transportation and storage):
Temperature	-25°C to +50°C (-13°F to +122°F)
Humidity	5% to 100% R.H., non-condensing
Air pressure	48kPa to 110kPa
Operating conditions	The system may not meet all performance specifications if operated outside of the following conditions:
Temperature	+18°C to +45°C (+59°F to +113°F)
Humidity	20% to 90% R.H. at +40°C, non-condensing
Air pressure	70kPa to 110kPa

