Safety of the medical gas pipeline system

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Abstract

Medical gases are nowadays being used for a number of diverse clinical applications and its piped delivery is a landmark achievement in the field of patient care. Patient safety is of paramount importance in the design, installation, commissioning, and operation of medical gas pipeline systems (MGPS). The system has to be operational round the clock, with practically zero downtime and its failure can be fatal if not restored at the earliest. There is a lack of awareness among the clinicians regarding the medico-legal aspect involved with the MGPS. It is a highly technical field; hence, an in-depth knowledge is a must to ensure safety with the system.

Keywords: Central supply, hospital, nitrous oxide, oxygen

Introduction

Gas pipeline failures have been reported several times in the anesthesia literature. Petty reviewed pipeline-related deaths from 1972 to 1993 in the US and reported 26 deaths which were due to crossed pipelines, defective connectors, and error in the supply. A survey by the Anesthesia Patient Safety Foundation showed a significant knowledge deficit among anesthesiologists regarding the medical gas pipeline systems (MGPS), and adequacy of knowledge is a must to make the system safe. The past decade witnessed many medico-legal cases compelling the clinicians to shift focus toward practical orientation of the system. Hence, maintaining safety standards is of prime importance starting from the main source at the manifold room to the final delivery point. The two main standards pertaining to MGPS are National Fire Protection Association 99 (US) and HTM 02-01 (UK). Others are British Standards EN 737 (BS EN 737), Compressed Gas Association, Canadian Standards Association, and the International Standards Organization. Cylinders should follow American Society for Testing and Materials standards, and pipes should have Lloyd’s certification as per BS 2871.

The core of the entire system is the manifold room which should be manned round the clock with trained personnel. It should have a suitable acoustic enclosure with 100% generator backup and fire security. The location should be marked clearly for ease of identification in the event of an emergency. It houses the control panel which allows a flow of 3000 L/min at 4.1 bar from the vacuum insulated evaporator (VIE) and relays alarm to secondary panels located throughout the hospital.

Oxygen

Continuous supply of oxygen is the primary requisite of any medical unit. According to BS EN 737-3:2000, there should be three independent supply sources. Primary, secondary, and a reserve source adequate to meet the demand in the event of primary and secondary supply failure. The manifold room should have 2 banks of D-type cylinders, each holding a minimum of 2 days consumption, attached to a fully automatic changeover control panel. Three-day consumption should be kept in reserve, as a contingency for Testing and Materials standards, and pipes should have Lloyd’s certification as per BS 2871.

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How to cite this article: Sarangi S, Babbar S, Taneja D. Safety of the medical gas pipeline system. J Anaesthesiol Clin Pharmacol 2018;34:99-102.
Aluminum or steel cylinders with Kevlar or Carbon fiber outer shell allow higher filling pressure, enabling larger storage. Cylinders should be tested by manufacturers at intervals of 5 years. Besides, oxygen concentrators or pressure swing adsorber system can supply a $\text{FiO}_2$ ranging from 0.95 to 0.97, strictly monitored by a paramagnetic oxygen analyzer. Liquid oxygen is an economical and convenient form of oxygen storage. The cryogenic liquid is stored in a VIE, usually of the capacity of 5 KL or 10 KL, although smaller Cryospeed™ vessels are also available. VIE should be installed in a high security and fire safe zone after taking regulatory approval from the Fires and Explosives Act office in Nagpur.

**Compressed Air**

It is used both as a driving force for equipment such as pneumatic drills (surgical air) or as an inhalational gas (medical air). The plant must ensure a flow of 3 KL/min at 8 bar, reduced thereafter as per requirement. Medical air needs a flow rate of 80 L/min at 4 bar and surgical airflow at the rate of 350 L/min at 7 bar. The medical air quality should meet the standards laid by the European Pharmacopoeia, restricting the carbon monoxide level to 5 ml/m³. Integral dryers, filters, and dew point monitor control the humidity to its allowable limit of 67 ml/m³.

**Vacuum and Anesthetic Gas Scavenging System**

Vacuum pressure of $-300$ mmHg is required at the terminal unit with a flow of 40 L/min.

Anesthetic gases are considered to be substances hazardous to health as per the Control of Substances Hazardous to Health Regulations 2002 except where they are administered to a patient in the course of treatment. Exhaust of both the systems should be carefully positioned away from the windows and intake of air compressors and ventilators. To control the greenhouse effect of the anesthetic gases, the anesthetic gas scavenging systems should incorporate a canister system which captures the unused gases, filters, and recycles them.

**Pipeline System**

Copper seamless pipes with fluxless silver brazing are used which should be as per ASTM standard and Lloyd’s certification. They are intercepted by the area valve service units (AVSUs) and area alarm panels (AAPs) [Figures 1 and 2]. AVSUs are placed in each clinical sector, to cutoff the gas delivery to the area beyond it during maintenance or to handle emergency. AAPs display the line pressures and have audiovisual alerts. All pipelines should be color coded with colored bands put at intervals of every 3 m [Figure 3].

**Outlet Points**

These are the final delivery points which are color coded, incorporating either the diameter index safety system or are of the quick connect type, available in two varieties, the Diamond and Chemetron [Figures 4 and 5]. The Chemetron is a more sturdy variant and is resistant to leakage.

Every new installation needs to be tested and verified as per the laid guidelines before putting the system into use. 

It is important to have a contingency plan to avoid crisis situations [Table 1].

| Testing Category | Description |
|------------------|-------------|
| Blowdown         | Lines are blown clear using oil-free dry nitrogen |
| Initial press test | System is subjected to 1.5 times working pressure to check leaks |
| Standing press test | System is subjected to 20% higher pressure for 24 h |
| Piping purge     | Purging of each outlet until there is no discoloration of the white cloth held over it |
| Cross-connection test | One gas system at a time using O2 analyzer |
| Final tie-in test | Active vacuum pipeline joints are tested using an ultrasonic leak detector |
Indigenous arrangement – all critical areas should have bulk oxygen cylinders, fitted with a double-stage regulator, tubing, and an adaptor. In case of manifold failure, the AVSU of the area is closed. The pipeline beyond it can now be fed with oxygen from this cylinder, by connecting the adaptor to any oxygen outlet point within the territory. Crisis due to vacuum pipeline failure can be tided over with portable electrical suction units.

The tender terms and conditions should specify that the supplier’s bulk oxygen trailer should reach the hospital as soon as possible in the eventuality of pipeline failure.

However, besides quality control and knowledge, safety can be ensured only when there is a proper upkeep and maintenance of the system. This can be ensured by:

- Formulating Standard Operating Procedures (SOP’s)
- Maintaining logbooks
- Keeping all the equipment under Annual Maintenance Contract/ Comprehensive Maintenance Contract (AMC/CAMC), allotted to the original supplier of the Equipment or to vendors with minimum 3-year experience in the field
- Preventive maintenance of equipment and leak test of pipeline should be ensured on quarterly basis
- 24 h manning by trained personnel
- Periodic training of manifold personnel
- Regular inspection of the pipelines, particularly after Public Works Department (PWD) work of every area
- Daily checking of contingency plan
- Mock drills of pipeline failure, fire, and explosion should be regularly conducted.

The vastness of the system has always remained a matter of concern whenever the issue of safety was raised. The anesthesiologists are an integral part of this system, but keeping in mind the medico-legal liabilities, the administrative control of the system should be vested upon the biomedical engineers, and it is imperative that all users should also be adequately trained.

Financial support and sponsorship
Nil.

Conflicts of interest
There are no conflicts of interest.
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