

Name of Policy: <u>General oxygen and aerosol equipment assembly</u> Policy Number: 3364-136-01-18 Department: Pulmonary Services Approving Officer: Senior Hospital Administrator Responsible Agent: Director, Pulmonary Services Scope: The University of Toledo Medical Center Pulmonary Services Department	 <p>Effective Date: June 1, 2023 Initial Effective Date: 12/1/2004</p>
<input type="checkbox"/> New policy proposal <input type="checkbox"/> Major revision of existing policy	
<input checked="" type="checkbox"/> Minor/technical revision of existing policy <input type="checkbox"/> Reaffirmation of existing policy	

(A) Policy Statement

All Respiratory Care practitioners and staff technicians dealing with the following equipment shall use the assembly methods described below.

(B) Purpose of Policy

To ensure safe set up and operation of the following equipment.

(C) Procedure

The following are instructions for the set up and use of equipment to be used by the Respiratory Care Department. Guidelines listed below are according to manufacturer suggestions as well as accepted standards of practice:

1. Cold Aerosol Oxygen Delivery:

Disposable aerosols are used for the administration of humidified oxygen ranging from oxygen concentration (FiO₂) of 21% to 100%.

- a. A flow top and its water return tube should be connected to the top of a pre-filled container of sterile water.
- b. The flow top with the container of sterile water should then be attached to the appropriate gas outlet, pressure regulator or N.C.G. medical gas outlet.
- c. The flow top should be adjusted to the prescribed FiO₂ and turned on with the flow of gas corresponding to the appropriate FiO₂ (between 5 and 10 liters per minute).
- d. Corrugated tubing is attached from the flow top to either a T- piece, trach collar, face mask or face tent.

2. The up-draft or hand-held nebulizer, used for the aerosolization of medications:

- a. The Up-Draft Nebulizer is a disposable plastic device driven by a medical gas source of either air or oxygen at 8 liters per minute, or manufacturer's recommended flow.
- b. The medical source gas is connected to the nebulizer and a T- piece for administration via mouthpiece, ventilator circuit, or directly from nebulizer to face mask or, trach collar. The patient should be monitored for pulse, respiratory rate, breath sounds, adverse reactions and general tolerance of treatment.

3. The ConchaTherm Neptune humidifier, used to humidify gas delivered by mechanical ventilators, and other situations where heated humidification is indicated:

Heated Wire mechanical ventilator system and High Flow oxygen system. The ConchaTherm system will be operated according to manufacturer’s recommendations.

Approved by:		Review/Revision Date:
<u>/s/</u>	<u>06/22/2023</u>	04/04/2001
Michael Taylor	Date	01/06/2005
Director, Pulmonary Services		03/05/2008
		03/09/2011
		08/03/2012
		12/01/2015
		04/08/2019
<u>/s/</u>	<u>08/01/2023</u>	12/30/2022
Shahnaz Rehman, M.D.	Date	06/01/2023
Medical Director		
<u>/s/</u>	<u>06/29/2023</u>	
Russell Smith	Date	
Senior Hospital Administrator		
<i>Review/Revision Completed By:</i> Director, Pulmonary Services		Next Review Date: June 1, 2026
Policies Superseded by This Policy:		

It is the responsibility of the reader to verify with the responsible agent that this is the most current version of the policy.