

Dr. Dry Saliva Absorbents Instructions for Use (IFU)

CAUTION:

Professional use only, ^R. These instructions, in whole or in part, are not a substitute for formal training. Appropriate professional education is REQUIRED prior to using this device clinically. Dr. Dry saliva absorbents are intended to be used by qualified dental practitioners in dental clinics, hospitals, labs, or schools for dental applications.

DESCRIPTION:

The Dr. Dry products are manufactured in two sizes, Large and Small. The Dr. Dry are manufactured with two backing types, Reflective, a metalized film backing, and Original, a white backing. The Dr. Dry are used to absorb saliva from the parotid gland. They have a semi-permeable barrier to capture the saliva with super absorbent particles and trap the saliva within the body of the Dr. Dry. Within the Dr. Dry is a core stiffener to offer cheek retraction as the product expands with saliva.


INDICATIONS:

Dr. Dry are indicated for anyone requiring dental care. Dr. Dry are intended to absorb moisture from the parotid gland for up to 15 minutes. The super absorbent polymer traps moisture as a gel when in contact with saliva. The Dr. Dry core stiffener provides cheek retraction to facilitate procedures far back in the oral cavity. The Dr. Dry backings brighten the oral cavity during procedures and the medical grade poly netting adhere gently to the tissue to stay in place yet removes easily with water spray.

CONTRAINDICATIONS TO USE:

Use of MDT's Dr. Dry is contraindicated on any patient who is allergic to any of the components of the product.

CLINICAL PRECAUTIONS AND WARNINGS:

- Do not use the product if package is opened or damaged. Discard any damaged Dr. Dry immediately.
- Do not use if the product is damaged.
- The Dr. Dry are single use. 
- To remove without tissue irritation, release adhesion with ample water spray to the buccal side of the Dr. Dry.
- Non-toxic by ingestion. If product bursts, remove as much as possible from mouth. Rinse mouth thoroughly with plenty of water. If adverse symptoms appear, seek medical attention.
- Always keep track of Lot Numbers or Dr. Dry to ensure traceability.

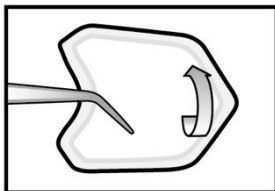
Failure to follow the instructions for use may lead to the following: allergic reaction, product leakage, procedural delay, minor irritation and discomfort, and cross-contamination.

CLINICAL USE:

Select the Dr. Dry size which adequately covers the buccal mucosa.

Step #1

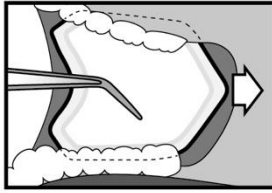
- Insert the Dr. Dry as shown with the absorbent pad against the cheek.



Step #2

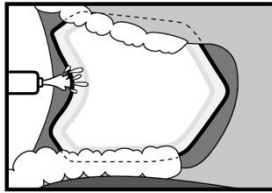
- Point the Dr. Dry to the back of the mouth as shown. In a few seconds, the Dr. Dry will begin to adhere to the

tissue and stay in place.



Step #3

- **Important:** To remove without irritating tissue, release adhesion with ample water spray to the buccal side of the Dr. Dry.



STORAGE:

Plantent Orbis Dry Pads should be stored in a dry, closed container. Improper storage conditions will shorten the shelf life and may cause malfunction of the product.

DISPOSAL

- Each used saliva absorbent must be disposed in a biohazards waste container.
- Each unused saliva absorbent may be disposed in a regular waste container.

PRODUCT SHELF LIFE

- Product shelf life is ten (10) years unless product is damaged.

TRACEABILITY:



Each package includes **Lot number** LOT on its label.













This number must be quoted in any correspondence regarding the product.

NOTICE: If a serious incident has occurred in relation to the device, the incident shall be reported to the manufacturer and if applicable, the competent authority of the Member State in which the user and/or patient is established.

To request a paper IFU free of charge, please contact Microcopy at sales@microcopydental.com or 800.235.1863, and an IFU will be delivered within seven (7) days.

SYMBOLS:

	Manufacture	Indicates the medical device manufacturer.		Do not reuse	Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.
---	-------------	--	---	--------------	--

	Lot Code	Indicates the manufacturer's batch code so that the batch or lot can be identified.		Consult instructions for use	Indicates the need for the user to consult the instructions for use.
	CE marking	Signifies European technical conformity.		Do not use if package is open or damaged	Indicates a medical device that should not be used if the package has been damaged or opened.
	Catalog Number	Indicates the manufacturer's catalog number so that the medical device can be identified.		Date of Manufacture	Symbol for date of manufacture.
	Medical Device	Indicates device is designed and intended for medical use.		Keep Dry	Indicates a medical device that needs to be protected from moisture.
	DEVICE for professional use only	(ref US FDA CDRH) Indicates device shall only be used by a trained professional.		Importer	Indicates the entity importing the medical device into the locale
	Distributor	Indicates the entity distributing the medical device into the locale		Authorized European representative	Indicates the Authorized representative in the European Community.

CONTACT INFORMATION:



Microcopy Dental
3120 Moon Station Rd. NW
Kennesaw, GA 30144, USA
sales@microcopydental.com
800.235.1863



MDT Micro Diamond
Technologies Ltd. 2
Ha'amal St. AFULA,
1857107, ISRAEL
www.mtdental.com



Obelis
Bd. Général Wahis 53
1030 Brussels, Belgium
Tel: + (32) 2. 732.59.54
Fax: +(32) 2.732.60.03
E-Mail: mail@obelis.net

REVISION HISTORY:

MCD-IFU-020 Rev. 2
Date of Issue: 03Feb2026