

Dade® Owren's Veronal Buffer

OV **BUFFER**



Revision bar indicates update to previous version.

Intended Use

Dilution buffer for coagulation testing.

Summary and Explanation

Owren's Veronal Buffer has been specifically prepared for coagulation studies but may be used in any laboratory test utilizing an isotonic barbital buffer, pH 7.35.

Reagents

| Reagent | Description | Storage | Stability |
|---------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------|------------------------------------|
| Dade® Owren's Veronal Buffer OV BUFFER | Ready to use liquid containing: <ul style="list-style-type: none"> • 2.84×10^{-2} M sodium barbital in 1.25×10^{-1} M sodium chloride pH 7.35 ± 0.1 | 2–8 °C May be used up to the expiry date indicated on the label if stored unopened. | 2–8 °C: once opened, 8 weeks |

Warnings and Precautions

For *in-vitro* diagnostic use only.

For laboratory professional use.

According to EU regulation 2017/746, any serious incident that has occurred in relation to the device shall be reported to the manufacturer and the competent authority of the EU Member State in which the user and/or patient is established.

Safety data sheets (MSDS/SDS) available on [siemens-healthineers.com/sds](https://www.siemens-healthineers.com/sds).

The pipetting steps must be performed with care in order to avoid contamination. Visible turbidity and flocculation are signs of microbial contamination.

Not for internal or external use by humans or animals.

Dispose of hazardous or biologically contaminated materials according to the practices of your institution. Discard all materials in a safe and acceptable manner and in compliance with all government requirements.

Preparing Reagents

OV **BUFFER** is ready for use.

More details on the use of **OV** **BUFFER** are given in the corresponding assay protocols or analyzer-specific Application Sheets provided by Siemens Healthineers.

Procedure

Materials Provided

| REF | Contents | | |
|----------|-------------------------------------------|------|-------|
| B4234-25 | Dade® Owren's Veronal Buffer OV BUFFER | 10 × | 15 mL |

Materials Required but not Provided

| Item | Description |
|--------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Coagulation analyzers ^a , such as: | <ul style="list-style-type: none"> • Atellica® COAG 360 System • BCS® XP System • SYSMEX CA-500/CA-600 series • SYSMEX CA-1500 System • SYSMEX CS-2000i/CS-2100i System • SYSMEX CS-2500 System • SYSMEX CS-5100 System • SYSMEX CN-3000/CN-6000 System |

^a Availability of analyzers may vary by country.
























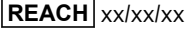
Please note that the applications on other analyzers can be validated by the instrument manufacturer in accordance with the requirements of the REGULATION (EU) 2017/746 under their responsibility as long as the intended purpose and performance are not modified.

Technical Assistance

For customer support, contact your local technical support provider or distributor.
siemens-healthineers.com

Definition of Symbols

The following symbols may appear on the product labeling:

| | | | |
|-------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------|-------------------------------------------|
|  | Do not reuse |  | Use By |
|  | Batch Code |  | Catalogue Number |
|  | Caution |  | Manufacturer |
|  | Authorized representative in the European Community |  | Contains sufficient for <n> tests |
|  | Biological Risks |  | <i>In Vitro</i> Diagnostic Medical Device |
|  | Temperature Limitation |  | Consult instruction for Use |
|  | Non-sterile |  | CE marking of conformity |
|  | CE marking of conformity with notified body ID number. Notified body ID number can vary. |  | Contents |
|  | Reconstitution volume |  | Level |
|  | Keep away from sunlight and heat |  | Warning |
|  | Danger |  | Prescription device (US only) |
|  | Device Identification (UDI) barcode |  | REACH Authorization Number |

Legal Information

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