



For the Analysis of Ethylene Glycol and Diethylene Glycol in Glycerin, Propylene Glycol, and Sorbitol via GC-FID (USP Monographs)

A turn-key solution, complete with chemicals, instruments, standard/sample preparation, analytical conditions, and software for operational and automated reporting.

- End-to-End workflow available
- Achieve precise determination of EG and DEG in pharmaceutical and food raw materials
- Simplifies instrument setup, analysis procedures, and compliance with regulatory requirements in QC labs
- Automatically generates test reports



Scan the QR code for more information on the Brevis GC

https://www.shimadzu.com.sg/an/pr oducts/gas-chromatography/brevis-g c.html





Complete SOP for the end-to-end workflow solutions

Easy to Use

Routine Analyzer

End-to-end workflow which comes complete with

- Standard Operating Procedure (SOP)
 - Sample preparation procedures
 - Software workflow for data analysis
 - Batch, method, and reporting templates
- Reagent list
- Consumables list

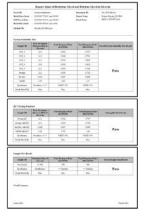
Validated Methods

According to USP Monographs

- For glycerin, propylene glycol, and sorbitol sample matrices
- **Pre-optimized analytical conditions** to ensure best performance
- Users only need to select the project accordingly to change conditions
- Any parameters can be changed, if necessary



Select the ready projects for each raw material with analytical parameters optimized according to the respective **USP Monographs**



Pre-set criteria for the automated calculation and reporting can be changed, if necessary

Automated Reporting

with Shimadzu MultiData Reporting (MDR)

- Checking of analytical results with automatic pass/fail judgement
 - System suitability test
 - QC closing standard
 - Tested sample final results based on USP monograph acceptance criteria
- Validation supported by Shimadzu Informatics team

Regulatory Compliance

with LabSolutions DB/CS

With an expansion of data integrity requirements around the world, the regulation is getting more stringent with each updated version. We continuously upgrade software security policies to keep up with current regulatory requirements.

- **Data protection and privacy** from unauthorized access
- **Records retention** from data loss with automatic backups
- Data integrity and authenticity in accordance to regulations such as FDA 21 CFR Part 11





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