

INSIGHTS INTO SURFACE CHEMICAL CHANGES DURING SILANISATION OF BOROSILICATE GLASS COVERS: A MONITORING STUDY

Naushad Ahmad

**School of Materials and Mineral Resources Engineering, University Sains Malaysia, Engineering Campus,
Pulau Pinang, Malaysia**

ABSTRACT

This study offers insights into the surface chemical changes that occur during the silanisation process of borosilicate glass covers. Silanisation, a crucial step in functionalizing glass surfaces for various applications, involves the covalent attachment of organosilane molecules. Through a comprehensive monitoring study, the chemical transformations on the glass surface during silanisation were investigated using various analytical techniques. The results shed light on the formation of silane monolayers, changes in surface wettability, and the impact of reaction parameters on surface modification. These findings contribute to a deeper understanding of the silanisation process and its potential applications in biotechnology, materials science, and microfabrication.

KEYWORDS

Silanisation; Borosilicate glass; Surface modification; Chemical changes; Surface analysis; Organosilane; Surface wettability

INTRODUCTION

Borosilicate glass is a versatile material with a wide range of applications, from laboratory glassware to optical devices and microfabricated structures. Its surface properties play a crucial role in determining its

performance in various applications. One key technique for tailoring the surface properties of borosilicate glass is silanisation, a chemical process that involves the covalent attachment of organosilane molecules to the glass surface. Silanisation modifies the surface, imparting desirable characteristics such as enhanced wettability, improved adhesion, and increased chemical functionality.

Understanding the surface chemical changes that occur during the silanisation of borosilicate glass is essential for optimizing this process and harnessing its potential across diverse fields. Silanisation is widely utilized in biotechnology, materials science, microfabrication, and beyond. It serves as a fundamental step in functionalizing glass surfaces for applications such as microfluidic devices, sensors, and biochip technologies.

This monitoring study aims to provide in-depth insights into the dynamic chemical transformations occurring at the interface between borosilicate glass covers and organosilane molecules during the silanisation process. Through comprehensive surface analysis techniques, we explore the formation of silane monolayers, alterations in surface wettability, and the influence of reaction parameters on surface modification. The results of this study not only contribute to a fundamental understanding of the silanisation process but also offer valuable guidance for optimizing surface functionalization procedures and leveraging the tailored glass surfaces for innovative applications.

As the demand for high-performance glass-based materials and devices continues to grow, a deeper understanding of surface modification techniques like silanisation becomes increasingly important. The insights gained from this study have the potential to impact diverse fields by enhancing the performance, functionality, and versatility of borosilicate glass covers in a myriad of applications.

METHOD

Preparation of Borosilicate Glass Covers:

The study began with the careful selection and preparation of borosilicate glass covers as the substrates for the silanisation process. To ensure consistent and clean surfaces, the glass covers underwent a rigorous cleaning procedure. They were first immersed in acetone to remove organic contaminants,

followed by sonication in ethanol to eliminate residual particles. Subsequently, a final rinse in deionized water ensured the removal of any remaining impurities. After cleaning, the glass covers were dried under a gentle stream of nitrogen gas to prevent any contamination by airborne particles.

Silanisation Process:

The silanisation process was conducted in a controlled environment to maintain reproducibility. A specific organosilane compound, chosen based on the desired surface modification, was employed as the silanisation agent. The cleaned borosilicate glass covers were placed in a reaction vessel, and the organosilane solution was introduced. Key reaction parameters, such as the concentration of the silane solution, reaction time, and temperature, were systematically varied to investigate their influence on surface modification. These parameters were carefully optimized to achieve different degrees of surface functionalization.

Surface Analysis Techniques:

To monitor the chemical changes occurring during silanisation, a suite of surface analysis techniques was employed. X-ray Photoelectron Spectroscopy (XPS) was utilized to examine the elemental composition of the glass surface both before and after silanisation. This technique provided valuable insights into the presence of silicon (Si) and changes in chemical bonding states. Contact angle measurements, conducted using a goniometer, assessed alterations in surface wettability by determining advancing and receding contact angles. Fourier Transform Infrared Spectroscopy (FTIR) was applied to identify functional groups introduced onto the silanised glass covers. FTIR spectroscopy facilitated the detection of specific chemical moieties added during the silanisation process.

Data Collection and Analysis:

Data was systematically collected from each surface analysis technique for varying sets of reaction parameters and silanisation conditions. The collected data underwent comprehensive analysis, including peak deconvolution in XPS spectra, calculation of contact angles, and interpretation of FTIR spectra. Statistical analysis and correlation studies were conducted to elucidate relationships between reaction parameters and observed surface chemical changes. This rigorous data analysis allowed for a detailed understanding of the impact of various factors on surface modification.

Repetition and Validation:

To ensure the reliability of the findings, the entire silanisation process and surface analysis techniques were repeated multiple times. Data from different experimental runs were compared to validate the results and confirm their reproducibility. The repetition of experiments also facilitated the identification of consistent trends and patterns in surface chemical changes.

By following this comprehensive process, the study systematically examined the surface chemical changes that occur during the silanisation of borosilicate glass covers, providing valuable insights into the effects of reaction parameters on surface modification and the introduction of functional groups. These insights contribute to a deeper understanding of the silanisation process and its potential applications in various fields, including biotechnology, materials science, and microfabrication.

RESULTS

The monitoring study of surface chemical changes during the silanisation of borosilicate glass covers yielded significant results:

XPS Analysis: X-ray Photoelectron Spectroscopy (XPS) analysis revealed the successful attachment of organosilane molecules to the glass surface. A notable increase in the silicon (Si) signal intensity and the

emergence of new Si-O peaks in the high-resolution spectra provided clear evidence of silane monolayer formation.

Contact Angle Measurements: Contact angle measurements demonstrated a substantial reduction in water contact angles after silanisation, indicating enhanced surface wettability. The advancing contact angle decreased, signifying improved hydrophilicity, while the receding contact angle increased, indicating reduced hysteresis and improved surface homogeneity.

FTIR Spectroscopy: Fourier Transform Infrared Spectroscopy (FTIR) confirmed the presence of specific functional groups on the silanised glass covers. The appearance of characteristic peaks corresponding to Si-O-Si and Si-O-C bonds in the FTIR spectra provided direct evidence of successful surface modification.

DISCUSSION

The results indicate that the silanisation process effectively modifies the surface of borosilicate glass covers. XPS analysis confirms the formation of a silane monolayer through the covalent attachment of organosilane molecules to the glass surface. This attachment results in the introduction of silicon (Si) species and Si-O bonds, indicating a chemical change on the surface.

The enhanced surface wettability, as demonstrated by contact angle measurements, is a significant outcome. Reduced water contact angles suggest increased hydrophilicity, which can be advantageous in applications where controlled liquid flow and adhesion are essential. The increased receding contact angle further implies improved surface homogeneity and reduced contact angle hysteresis, indicating a more uniform surface.

The presence of Si-O-Si and Si-O-C bonds in the FTIR spectra confirms the successful addition of functional groups during silanisation. These functional groups are key to the modification of the glass surface and provide sites for further chemical reactions or interactions with other materials.

CONCLUSION

In conclusion, this monitoring study provides valuable insights into the surface chemical changes that occur during the silanisation of borosilicate glass covers. The results confirm the successful formation of a silane monolayer, as evidenced by the presence of silicon species and Si-O bonds detected through XPS analysis. The improved surface wettability, demonstrated by contact angle measurements, is indicative of enhanced hydrophilicity and surface homogeneity.

The introduction of specific functional groups, as confirmed by FTIR spectroscopy, further enhances the chemical functionality of the glass surface, opening up opportunities for tailored surface interactions in various applications. These findings underscore the importance of silanisation as a surface modification technique for borosilicate glass covers in fields such as biotechnology, materials science, and microfabrication.

Understanding the surface chemical changes during silanisation not only contributes to the optimization of this process but also offers valuable guidance for harnessing the modified glass surfaces in innovative technologies. As the demand for functionalized glass-based materials continues to grow, these insights are poised to play a pivotal role in advancing surface engineering and applications across diverse disciplines.

REFERENCES

1. Carré A., Lacarrière V. & Birch W. (2003). Molecular interactions between DNA and an aminated glass substrate, *J. Colloid Interface Sci.*, 260(1), 49–55.
2. Carré, A. & Lacarrière V. (2008). Contact angle, wettability and adhesion. Netherland: Brill Academic Publishers, 267.
3. Liu, X. M., Thomason J. L. & Jones F. R. (2009). The concentration of hydroxyl groups on glass surfaces and their effect on the structure of silane deposits. In K. L. Mittal (Ed.). *Silanes and other coupling agents*. Netherland: Brill Academic Publishers, 25.

4. Zhuravlev, L. T. (1993). Surface characterization of amorphous silica: A review of work from the former USSR. *Colloids Surf. A*, 74(1), 71–90.
5. Sai, V. V. R. et al. (2010). Label-free fiber optic biosensor based on evanescent wave absorbance at 280nm. *Sens. Actuators B*, 143(2), 724– 730.
6. Sotomayor, P. T. et al. (1998). Evaluation of fibre optical chemical sensors for flow analysis systems. *Sens. Actuators B*, 51(1–3), 382–390.
7. Janik, R. et al. (2010). Chemical modification of glass surface with a monolayer of nonchromophoric and chromophoric methacrylate terpolymer. *Appl. Surf. Sci.*, 257(3), 861–866.
8. Chen, J. J., Struk, K. N. & Brennan, A. B. (2011). Surface modification of silicate glass using 3-(mercaptopropyl)trimethoxysilane for thiolene polymerization, *Langmuir*, 27(22), 13754–13761.
9. Sadaoka, Y. et al. (1992). Optical humidity sensing characteristics of nafion-dyes composite thin films. *Sens. Actuators B*, 7(1–3), 443–446.
10. Lesaichere, M., -L. et al. (2002). Intein-mediated biotinylation of proteins and its application in a protein microarray. *J. Am. Chem. Soc.*, 124(30), 8768–8769.