Lab Report: Analysis of Natural Oil Mixtures

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Introduction

This report documents the analysis of various natural oil mixtures using advanced instrumentation. The samples include

combinations of oils and additives commonly found in cosmetic and pharmaceutical applications. Each mixture

underwent a series of tests to investigate its physical and chemical properties.

Methodology and Observations

The following instruments were employed to assess the characteristics of each sample:

1. Titrator T-905: Utilized for determining acidity or basicity levels.

2.UV-Vis Spectrophotometer UV-2600: Employed to measure absorbance and transmittance.

3.X-Ray Diffractometer XRD-6000: Used for analyzing crystal structures.

4.PCR Machine PCR-96: Applied for amplification of nucleic acids, though in this context, Ct values are recorded.

5. Microplate Reader MRX: Measures optical density at various wavelengths.

6.HPLC System HPLC-9000: Conducted high-pressure liquid chromatography for compound separation.

7.Gas Chromatograph GC-2010: Utilized for separating and analyzing compounds that can be vaporized.

8. Rheometer R-4500: Used to assess the flow properties of the mixtures.

9. Viscometer VS-300: Evaluated the viscosity of fluid samples.

Observations

During experimentation, varied texture and viscosity were observed across different samples. For instance, samples

containing beeswax presented higher viscosity, while those with cetyl alcohol demonstrated smoother consistency.

Vitamin E-rich samples showed increased stability.

Sample Measurement Data

The following tables outline the measurement readings obtained from each instrument:

Table 1 - Sample Identification and Procedure Results

	Instrument	Sample Composition	Measure Type	Result
	Titrator T-905	Coconut Oil, Beeswax	М	4.672
	-Vis Spectrophotometer UV-26	യ്യാമ Oil, Cetyl Alcohol, Glycer	in Abs	2.987
	(-Ray Diffractometer XRD-600	onut Oil, Cetyl Alcohol, Vitami	n E C	85.0

Table 2 - Chromatography and Miscellaneous Data

Instrument	Sample Composition	Units	Measurement
HPLC System HPLC-9000	Coconut Oil, Beeswax	mg/L	350.7
Gas Chromatograph GC-2011	joba Oil, Cetyl Alcohol, Glycer	in ppm	308.6
Rheometer R-4500 Co	conut Oil, Cetyl Alcohol, Vitami	n E Pa-s	875.4
Microplate Reader MRX AI	mond Oil, Cetyl Alcohol, Glyce	rin OD	3.8

Table 3 - Viscosity Measurements

Instrument	Sample Composition	Units	Viscosity
Viscometer VS-300	Jojoba Oil	сР	2459.32
Viscometer VS-300 (	oconut Oil, Beeswax, Vitamin	E cP	4849.9

Results and Discussion

The titration results suggest a moderate acidity in the coconut oil and beeswax composition, with a notable molarity of 4.672. Absorbance levels, as indicated by the UV-Vis Spectrophotometry, were highest in the sample containing Jojoba Oil and additives, which implies strong light absorption capabilities.

X-ray diffraction confirmed the crystallinity of the coconut oil, cetyl alcohol, and vitamin E mixture, with a confidence level (C) of 85. Interestingly, HPLC analysis revealed substantial concentrations of active compounds in the coconut oil and beeswax mixture, reflected in a high mg/L reading of 350.7.

Random Note: There was unexpected interference in the chromatography baseline during analysis, attributable to unrelated external equipment malfunctions. This did not affect the integrity of the results as it was accounted for in data processing.

PCR readings, particularly the Ct values for the Jojoba Oil, Gum, and Vitamin E, averaged around 32, suggesting potential interaction at a genetic analysis level, which would not generally be applicable in a non-biological context.

The gas chromatography data points to significant ppm concentrations in the Jojoba Oil mixture, essential insight for quality control and formulation sciences.

Rheometric analysis provided comprehensive shear stress data for the mixtures, especially coconut oil blends, indicative of complex viscoelastic behavior, which exceeded simple liquid flow behavior.

## Conclusion

Comprehensive analysis of various oil and additive mixtures highlighted noteworthy differences in physical and chemical properties. These findings underpin formulation development for applications where such bases are critical. Future investigations might explore thermal behavior and long-term stability of such mixtures under varying environmental conditions.

Appendix: Various other data points were recorded during the experimentation, some of which were later determined to be irrelevant to the current analysis scope, such as temperature fluctuations in lab conditions and experimental errors corrected during calibration checks.