

Application Note

DIAMAXATR™ DIAMOND ATR

NO. 21165



Figure 1. The DiaMaxATR.

Dissolution of Nail Polish Using Acetone

INTRODUCTION

Diamond ATR can be useful for nail polish or paint manufacturers for quality control as well as developing or altering formulations. This can also apply to paint thinner or stain remover manufacturers in testing the effectiveness of their products. In this study, the dissolution of nail polish with acetone is examined using the DiaMaxATRTM accessory.

EXPERIMENTAL

Infrared spectra were collected on an FT-IR spectrometer equipped with the Harrick DiaMaxATRTM single-reflection high throughput diamond ATR accessory. Spectra were collected at an 8 cm⁻¹ resolution, a gain of 1, and signal averaged over 32 scans. The aperture was set to 100%. Spectra were collected in the range 4000-400 cm⁻¹. The instrument was purged to reduce water vapor and CO_2 interferences.

The samples used were nail polish (O.P.I in "Over the Taupe") and acetone. The ingredients listed by O.P.I in their MSDS include solvents: ethyl acetate, butyl acetate, isopropyl alcohol and propyl acetate; plasticizers: camphor, trimethyl pentanyl diisobutyrate and ethyl tosylamide; and polymers: nitrocellulose and tosylamide/formaldehyde resin.

One coat of nail polish was applied on the diamond ATR crystal and allowed to dry completely for 20 minutes while spectra were collected every 30 seconds using Harrick's TempLink software. After a spectrum was taken of the dried nail polish, two drops of acetone were deposited on top of the dried nail polish. A sealed liquid cell was used to enclose the samples to reduce evaporation of acetone. Spectra were collected every 30 seconds for a total of 10 minutes.

RESULTS AND DISCUSSION

Figure 2 displays the spectra of nail polish before and after it dries for 20 minutes. Initially, there is a broad band between 3700-3200 cm⁻¹, which may be attributed to O-H bonds from the isopropyl alcohol. There are three peaks near 3000 cm⁻¹ (2966 cm⁻¹, 2937 cm⁻¹, and 2876 cm⁻¹), indicating there are C-H bonds, which could be from numerous ingredients, which include the polymer, solvents, resins, and plasti-

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ANALYSIS OF NAIL POLISH DISSOLUTION USING THE DIAMAXATR™

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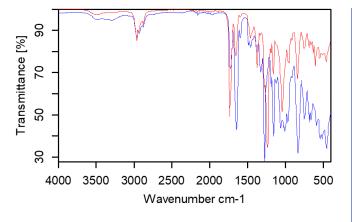


Figure 2. Spectra of nail polish right after 1 coat was applied on the diamond (red) and after the same coat of nail polish dried for 20 minutes (blue).

cizers. There is a peak at 1736 cm⁻¹ which is due to carbonyl (C=O) bonds found in the solvents: ethyl acetate, butyl acetate, and propyl acetate as well as one of the plasticizers, trimethyl pentanyl diisobutyrate. At 1656 cm⁻¹ there is a peak, which may be due to C=C stretching from cyclic alkenes in tosylamide/formaldehyde resin. After the polish dries, the initial band between 3700-3200 cm⁻¹ broadens and a second band emerges at 3300 cm⁻¹. Although it is likely the isopropyl alcohol has evaporated, N-H bonds from tosylamide/formaldehyde resin may have contributed to the bands between 3700-3200 cm⁻¹ as the solvents evaporate and concentrate the remaining ingredients. The evaporation of the solvents is shown more clearly when observing the (C=O) peak at 1736 cm⁻¹ because it decreases in intensity and shifts to 1725 cm⁻¹ as the nail polish dries. The (C=C) peak at 1656 cm⁻¹ increases in intensity and shifts to 1643 cm⁻¹ as the tosylamide/ formaldehyde resin concentrates.

Figure 3 compares a spectrum of nail polish immediately after two drops of acetone was deposited on the dry coat of nail polish and a spectrum collected 10 minutes later. Since the attraction between the polymer molecules are weaker than the attraction

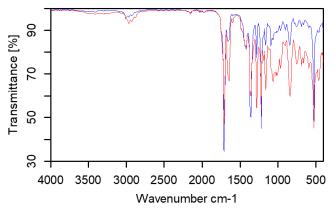


Figure 3. A comparison of a spectrum of dried nail polish right after 2 drops of acetone was deposited on top of it (red) and a spectrum collected 10 minutes later (blue).

between the acetone molecules and the polish polymer molecules, the polymer turns from solid to liquid. In figure 3, this is observed when comparing the (C=O) peak at 1710 cm⁻¹ after acetone was initially deposited and after 10 minutes. After 10 minutes with acetone, the peak intensity increased. As expected, the (C=C) peak at 1644 cm⁻¹ decreased in intensity and shifted to 1657 cm⁻¹.

CONCLUSION

Using diamond ATR, nail polish manufacturers can monitor the changes in their formulas as it dries to study how the effects of altering the formulation impact the overall performance of their products. Additionally, the dissolution of nail polish can also be examined to either enhance the durability of the polish or aid in the ease of removal. Similarly, paint thinner manufacturers can observe how their products perform over time while dissolving paint.

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