The photochemical reactions of methanolic nalidixic acid (NAL) solution in the absence and in the presence of air have been investigated using 300 nm UV light. From the reactions, 1-ethyl-7-methyl-4-oxo-4-hydro-1,8-naphthyridine (EMDN), formic acid, and formaldehyde are produced. In the presence of air, hydrogen peroxide is also detected along with the products listed above. The presence of oxygen during the irradiation of methanolic NAL solution effects on the product yield. The initial quantum yields of the products and of the NAL decomposition are determined. Possible reaction pathways for the photochemical reaction are suggested on the basis of the products analysis.

**Key Words:** Photochemical reaction, Nalidixic acid, 1-Ethyl-7-methyl-4-oxo-1,4-hydro-1,8-naphthyridine, Formic acid, Formaldehyde

**Introduction**

Quinolone derivatives are the most frequently used antibacterial reagents. Nalidixic acid (NAL) is one of quinolone derivatives and was synthesized by Lesher et al. in 1962.1 A number of quinolone derivatives have been synthesized since then. However, it is reported that these drugs are able to induce photosensitive reactions in human skin by sunlight.2,3 It is, therefore, interesting to study the photochemical reaction of the quinolone antibacterial reagents since it could directly present the photoproducts during their UV-irradiation. However, their photochemical properties were poorly investigated, and only a few papers were reported on the study of the photochemical reaction of NAL.

Dezer et al.4 reported that photolysis of oxygen-free nalidixic acid in the basic solution results in the loss of the COOH group at the 3-position. From the decarboxylation, 1-ethyl-7-methyl-4-oxo-4-hydro-1,8-naphthyridine (EMDN) is yielded. Carbon dioxide, ethylamine and 1-ethyl-1,4-dihydro-7-methyl-2,4-dioxo-1,8-naphthyridine are also produced using a polychromatic UV light. Douglas et al. tested the photosensitization with nalidixic acid and oxolinic acids as photosensitizers in aqueous solution using 365 nm light.5 Vermeersch et al. reported on the study of a laser flash photolysis.6 The 335 nm laser flash photolysis of nalidixic acid at pH 9.2 leads to the triplet state formation of the nalidixate anion. In 1990, Fernandez et al. proposed the mechanism of photo-photolysis of EMDN.7 It is, however, hardly reported not only on the quantitative investigation on the photochemical reaction pathway of NAL but also on the photolysis of NAL in nonaqueous solution. NAL can be dissociated in aqueous solution and we reported its pKa value was found to be 6.33.8

It is conventional to postulate photochemical reaction pathway based on the product analysis because photochemical reaction undergoes very fast and various products can be formed. The purpose of this study is to elucidate the photochemical reaction pathway of methanolic NAL in the absence and in the presence of air using the UV light of 300 nm based on the products analysis.

**Experimental Section**

**Reagents and General.** Nalidixic acid (NAL) was purchased from Sigma Chemical Co. (St. Louis, U.S.A.) and was used without further purification. The other chemicals were reagent grade and used as received. Methanol was first saturated by bubbling for 60 minutes with high purity argon and then 1 mM deaerated methanolic NAL solution was prepared. 80 mL of the prepared solution (1 mM) was transferred into the irradiation vessel and irradiated at room temperature using a photoreactor (Rayonet, RPR-208) equipped with a UV lamp as a monochromatic light source (λ = 300 nm).

The intensity of the lamp irradiance was measured by potassium ferrioxalate actinometry.9-10 The reduction yield of Fe³⁺aq to Fe²⁺aq ion after irradiation of 6 mM deaerated aqueous potassium ferrioxalate solution was measured by spectrophotometric method.11 The lamp intensity was found to be 7.66 × 10¹⁸ hv-L⁻¹-min⁻¹ from the known quantum yield, Q (Fe²⁺aq) = 1.24.9,10 No change in the intensity of the lamp irradiance was observed over the period of the experiment.

Steady-state fluorescence spectra of the 0.05 mM deaerated NAL in methanol-water mixtures were obtained with a spectrofluorometer (Perkin-Elmer, LS-50) at 25 °C. Methanol-water mixtures were prepared using doubly distilled water, which was obtained by passing distilled water through a deionization system (Barnstead, Nonopure II). Fluorescence quantum yields were determined using quinine sulfate as a reference.12-14 The absorption spectra were taken using a UV-Vis spectrophotometer (Uvikon, model 943, Italy).
Problems Analysis. To perform the qualitative analysis of the products after irradiation of NAL in methanol, 60 mL of the irradiated solution was concentrated to 5 mL using a rotary vacuum evaporator. The sample prepared was then analyzed by two kinds of analytical methods. One of them was using a GC-MS system (Varian saturn; DB-5 capillary column 60 m × 0.32 m, EI method); [product; m/z (rel. intensity), formaldehyde; 28(10), 29(100), 30(35), formic acid; 17(20), 29(100), 44(40), 46(60)]. Identifications and determinations were made by comparison with fragmentation patterns of known amounts of the pure substances. The other was analyzed using a LC-MS system (Mariner MS-spectrometer equipped with ESI interface, capillary C18 reversed column, eluent; 1 : 4 volume ratio of acetonitrile : water). Two MS spectra were obtained: one has 232 molecular weight indicating NAL and the other has 188 molecular weight. These were separated by column chromatography (eluent was 1 : 1 volume ratio of benzene:ethylacetate). The solution was then washed by methanol and dried. The $^1$H-NMR spectrum of the sample was obtained by a $^1$H-NMR spectrometer (Varian Gemini; CDC13, 300 MHz) $\delta$ 1.44 (3H, t, J = 7.2 Hz), 2.44 (3H, s), 4.40 (2H, q, J = 7.2 Hz), 6.30 (1H, d, J = 7.8 Hz), 7.19 (1H, d, J = 8.1 Hz), 7.66 (1H, d, J = 7.8 Hz), 8.58 (1H, d, J = 8.1 Hz) and it was characterized to be 1-ethyl-7-methyl-4-oxo-1,4-hydro-1,8-naphthyridine (EMDN).

Quantitative analysis of the products was carried out either by the liquid chromatographic method or by the spectrophotometric method because the gas chromatographic method has a poor reproducibility. Quantitative analysis of EMDN produced and NAL decomposed performed by a liquid chromatography (Young-Lin, Model 930; phase sep S5 ODS2 column, eluent; 30 vol.% of 2 mM aqueous KH2PO4 solution, UV detector) with a calibration curve using S5 ODS2 column, eluent; 30 vol.% of 2 mM aqueous KH2PO4 solution, UV detector) using citric pyridine as an internal standard. On the other hand, the amount of formic acid was determined by the same liquid chromatography equipped with a Ion-Exclusion column (IC-PAKTM; eluent: 0.01 N H2SO4, UV detector) using citric acid as an internal standard. The amount of formaldehyde produced was determined by spectrophotometric method. An aliquot (5 mL) of the irradiated solutions treated with Hantz reagent. The molar extinction coefficient ($\varepsilon$) of colored complex was 7860 M$^{-1}$cm$^{-1}$ at 412 nm in this experiment and it was not interfered by the presence of the other compounds such as methanol. Spectrophotometric determination of hydrogen peroxide was performed by treatment with acidic TiCl4 solution. The developed color was characterized by the absorbance maximum at 414 nm and its molar extinction coefficient ($\varepsilon$) at this wavelength was determined to be 3430 M$^{-1}$cm$^{-1}$.

Results and Discussion

Irradiation of 1.0 mM NAL in methanol was carried out using 300 nm UV light in the absence and in the presence of air. To examine the photochemical reaction, UV-Vis absorption spectra and pH values were measured before and after irradiation of methanolic NAL solution. As shown in Figure 1, their absorption spectra appeared as a similar shape in the wavelength region above 260 nm whereas they were different in the wavelength region below 260 nm. The absorption spectrum of aqueous NAL solution contains two bands in the wavelength region above 260 nm, indicating two chromophores: one assigned from the nitrogen atom at the position 1 to carboxyl group and the other from the methyl group attached to the 7-carbon atom to carbonyl group. The fact that the absorption spectra appeared as a similar shape in the wavelength region above 260 nm may imply that aromatic moiety in NAL was not so greatly altered after irradiation of NAL using 300 nm UV light. The pH value of the solution decreased from 6.59 to 5.43 when 6.89 × 10$^{20}$ hυ·mL$^{-1}$ UV dose were irradiated in the deaerated methanolic NAL solution. The same pH change occurred in the irradiation of aerated methanolic NAL solution, but its difference (0.71) was rather small compared to that of deaerated solution. It indicates that some acids were produced in the irradiation of methanolic NAL solution and oxygen affects the photochemical reaction process. After irradiation of the solution, 1-ethyl-7-methyl-4-oxo-4-hydro-1,8-naphthyridine (EMDN), formic acid and formaldehyde were mainly produced. The pH change of the solution before and after irradiation could be explained by the formation of formic acid.

The photolysis of 1.0 mM deaerated as well as aerated methanolic NAL solution was studied as a function of the number of quanta. As shown in Figure 2, the decomposition of NAL in methanol decreased but not in proportion to the number of quanta. The electronically excited NAL is not so much decomposed and it has some photostability in the initial state of the photochemical reaction. However, the decomposition of NAL increased regardless of the absence and the presence of air, as the number of quanta increased. This behavior indicates that some secondary photochemical reactions, which contribute to the decomposition of NAL, occurred in the system by the absorption of the UV light. In addition to this, the amounts of photoproducts such as EMDN, formic acid, and formaldehyde did not increased in proportion to the number of quanta as presented in Figure 3.
Therefore, to avoid the interference of secondary reactions, we determined the initial quantum yields (Q_i), obtained from the slope of tangent line of the curve shown in Figure 3. The results are summarized in Table 1.

In the irradiation of NAL in methanol by 300 nm UV light, methanol does not absorb the light and only NAL absorbs all of the light at the initial process of the reaction. It means that the photochemical decomposition of NAL in methanol begins with absorption of the light by NAL. The fact that EMDN and formic acid are mainly produced in the irradiation of the solution, is an evidence that the electronically excited NAL by the absorption of 300 nm is predominantly decomposed by splitting into the radical I and OOH as in Scheme 1.

The radicals of I and OOH produced can react with the chemical species in the solution. However, most of the radicals attack methanol because methanol has the largest concentration among the chemical species in the solution, and its reaction probability of the reactions defined by multiplication of concentration and rate constant is greatest. Hence, the formation of EMDN and formic acid could be explained by the reactions (1) and (2), respectively.

\[
\text{COOH} + \text{CH}_3\text{OH} \rightarrow \text{CH}_2\text{OH} + \text{HCOOH} \quad (2)
\]

The initial quantum yield of EMDN and that of formic acid produced by the irradiation of deaerated methanolic NAL were found to be nearly same. It supports that the hypothesis described above is reasonable because the products were formed by the radicals of I and COOH, and the amount of each radical produced by the photofragmentation of NAL was same. The photochemical reaction mechanism for the formation of EMDN is quite similar to that proposed by Dezer et al. \cite{4}, although they reported on the photolysis of oxygen-free nalidixic acid in the basic solution leading to the formation of the dissociated chemical species of NAL, so called nalidixate anion. However, NAL is hardly dissociated in methanol. It means that the photolysis of NAL in methanol was proceeds from its molecular state rather than from its anionic form. This is the reason why formic acid

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**Table 1.** Initial quantum yield (Q_i) of the compounds after irradiation (\( \lambda = 300 \) nm) of 1 mM nalidixic acid (NAL) in aerated and deaerated methanol

<table>
<thead>
<tr>
<th>Compounds</th>
<th>Q_i when saturated with Ar</th>
<th>Q_i when saturated with Air</th>
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<tr>
<td>NAL</td>
<td>( 5.51 \times 10^{-5} )</td>
<td>( 2.36 \times 10^{-5} )</td>
</tr>
<tr>
<td>EMDN</td>
<td>( 2.47 \times 10^{-5} )</td>
<td>( 1.46 \times 10^{-5} )</td>
</tr>
<tr>
<td>Formic acid</td>
<td>( 2.53 \times 10^{-5} )</td>
<td>( 4.19 \times 10^{-6} )</td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>( 2.31 \times 10^{-5} )</td>
<td>( 2.22 \times 10^{-5} )</td>
</tr>
<tr>
<td>Hydrogen peroxide</td>
<td>( - )</td>
<td>( \leq 10^{-6} )</td>
</tr>
</tbody>
</table>

*These values indicate the initial quantum yield of nalidixic acid decomposed. The others indicate the initial quantum yield of the products.*
was formed by the irradiation of NAL in methanol.

In general, the electronically excited state of aromatic compounds possesses larger dipole moment ($\mu'$) than that of the ground state ($\mu$). Therefore, it might be expected that the electronically excited NAL in methanol could be dissociated to nalidixate anion form more easily. To ascertain this expectation, we measured the spectroscopic properties such as Stokes shift and fluorescence quantum yield of NAL in the methanol-water mixtures. As shown in Table 2, Stokes shift of NAL decreased as the polarity of the mixtures decreased. This tendency is very general because the emission spectrum shifts to shorter wavelengths as the solvent polarity decreases. However, the fluorescence quantum yields of NAL were independent in the methanol-water mixtures except for that in water ($Q_F = 0.0054$). It indicates that the chemical structure of the electronically excited NAL in water was different from that in mixtures with methanol. If NAL could be dissociated to nalidixate anion form, the amount of fluorophore should be changed by the polarity of the mixtures and fluorescence quantum yield of NAL should be also changed. But, it was found to be nearly same value. This result could be interpreted by the formation of intramolecular hydrogen bond between the proton of the carboxyl group and the keto oxygen in NAL. This is a reason why carboxyl group in NAL was not dissociated in the electronically excited state, and why COOH radical could be produced by the irradiation of NAL in methanol.

The $\text{CH}_2\text{OH}$ radical formed by the processes (1) and (2) can predominantly attack methanol. Sonntag reported that hydrogen atom in methanol is mainly dissociated by attack of a radical.\textsuperscript{17} It is therefore expected that $\text{CH}_2\text{OH}$ radical is reproduced by the reaction of a $\text{CH}_2\text{OH}$ radical and methanol. Hence, the $\text{CH}_2\text{OH}$ radicals should be stabilized by the another reaction pathway such as the combination with each other to produce formaldehyde and ethylene glycol as in reaction (3). Therefore, the formation of formaldehyde could be interpreted by the reaction (3a).

$$\text{HCHO} + \text{CH}_2\text{OH} \rightarrow \text{CH}_2\text{OH}_2 \quad \text{(3a)}$$

$\text{CH}_2\text{OH}$ radicals produced in reaction (5) takes part in the formation of formaldehyde again. Therefore, the initial quantum yield of formaldehyde is nearly same value regardless of existence of air. To test this expectation, the amount of hydrogen peroxide formed by the irradiation of aerated NAL in methanol was analyzed. The formation of hydrogen peroxide produced by the irradiation of aerated NAL in methanol was detected. But its amount was so small. It is because the formation of $\text{H}_2\text{O}_2$ radical is also small. In contrast to the results, its formation was not detected in the irradiation of deaerated methanolic NAL.

**Conclusions**

1-Ethyl-7-methyl-4-oxo-1,8-naphthyridine (EMDN), formic acid, and formaldehyde were produced as a main product in the irradiation of NAL in methanol. The electronically excited NAL in methanol by the absorption of 300 nm UV light exists its molecular form rather than its dissociated form, and it is mainly split into its decarboxy-

<table>
<thead>
<tr>
<th>Vol. % of H$_2$O</th>
<th>$\nu_i$ (cm$^{-1}$)*</th>
<th>$\nu_f$ (cm$^{-1}$)*</th>
<th>$\Delta\nu$ (cm$^{-1}$)*</th>
<th>$Q_F$</th>
<th>$\frac{\nu_f}{\nu_i}$</th>
<th>$a$ (cm$^{-1}$)</th>
<th>$b$ (cm$^{-1}$)</th>
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<td>300</td>
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*Energy of absorption ($\nu_i$) or fluorescence center of gravity ($\nu_f$): uncertainty is ±60 cm$^{-1}$. \(\frac{\nu_f}{\nu_i}\): uncertainty of $\Delta\nu$ is ±6%.
ylated radical and $\cdot$OOH radical. The both radicals attack methanol to form EMDN and formic acid. Thus $\cdot$CH$_2$OH radical is produced and the formation of formaldehyde could be explained by the disproportionation process of the radical. The photochemical reaction of NAL was affected by the presence of oxygen in the solution. The decomposition of NAL as well as the formation of EMDN and formic acid better proceeded in the absence of air than in the presence of air. However, the amount of formaldehyde produced is not affected by the presence of air because $\cdot$HO$_2$ radical formed by the reaction between $\cdot$OOH radical and oxygen contributed to the formation of $\cdot$CH$_2$OH radical.

References