

### Physicochemical Properties of Chromium-doped Titanium Dioxide Mesoporous and Its Application for Antifogging Materials

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#### **ABSTRACT**

Mesoporous materials of chromium doped titanium dioxide (Cr-doped TiO<sub>2</sub>) and undoped TiO<sub>2</sub> were prepared by hot-injection reflux technique at 150 °C for 6 hours. Samples Cr-doped TiO, at different percentages: 1.1, 3.9 and 4.4 (wt% Cr) and undoped TiO, were synthesized from Ti(O<sub>2</sub>)O.2H<sub>2</sub>O as titanium source obtained from the reaction of TiCl<sub>4</sub> and H<sub>2</sub>O<sub>2</sub>. Solid (NH<sub>4</sub>)<sub>2</sub>CrO<sub>4</sub> was a source of chromium as dopant. The prepared materials were characterized using powder X-ray diffraction (PXRD), scanning electron microscopy-energy dispersive X-ray spectroscopy (SEM-EDS) and N, adsorption-desorption isotherm. The XRD results reveal that the undoped TiO2 is composed of well-crystalline anatase (major), rutile (minor) and brookite (minor) phases. In the 1.1 wt% Cr-doped TiO, its phase composition is anatase (major) and rutile (minor). The chromium dioxide (CrO<sub>2</sub>), anatase (major), brookite and srilankite (TiO<sub>2</sub>-II) are present in the 3.9 wt% Cr-doped TiO<sub>2</sub> and the 4.4 wt% Cr-doped TiO<sub>2</sub>. All prepared materials (Cr-doped TiO<sub>2</sub> and undoped TiO<sub>3</sub>) exhibit mesoporous of type-IV isotherm curves with H2-type hysteresis loop according to the IUPAC classication. The Brunauer-Emmett-Teller (BET) specific surface area ( $S_{BET}$ ) and the mean pore size of the 4.4 wt% Cr-doped TiO2 exhibit a maximum surface area of 111 m<sup>2</sup>/g, corresponding to mean porous size of 4.95 nm. The hydrophilic properties of Cr-doped TiO<sub>2</sub> were investigated with illumination of UV light. All prepared samples shows excellent superhydrophilic properties. The 4.4 wt% Cr-doped TiO, demonstrates the most excellent superhydrophilic properties as compared with the other samples. These results allow the materials to be prospective application as antifogging.

Keywords: titanium dioxide, Cr-doped TiO,, mesoporous, superhydrophilic, antifogging

#### 1. INTRODUCTION

Among the various semiconductors, titanium dioxide (TiO<sub>2</sub>) has been well known as an efficient phtocatalyst. This is because TiO<sub>2</sub> has the most efficient photoactivity, high refractive index, light absorption, non-toxicity, high chemical stability and relatively low-cost production [1]. When irradiated with ultraviolet or sun light on a TiO, surface, two phenomena of photochemical reaction will happen: the first is the photo-induced redox reactions, and the other is the photoinduced super-hydrophilic conversion. When the surface of TiO, was irradiated with light consisting of wavelengths shorter than its band gap, about 3.0-3.2 eV, electron and hole pairs are generated in the TiO<sub>2</sub>, and they reduce and oxidize adsorbates on the surface, generating radical species such as XO2 and ×OH. Super-hydrophilic surfaces and reduction reactions at the surface of TiO, are a broad research field covering such as water cleaning [2], photo-electrochemical splitting of water [3], solar cells [4], self-cleaning [5], antifogging [6], anti-bacterial surface coatings [7], and photocatalyst [8]. Various applications of self-cleaning TiO, films have been proposed especially for practical applications such as window glasses, mirrors and windshields of automobile [9].

The performances and the properties of TiO<sub>2</sub> are strongly influenced by crystalline structure, morphology, surface states, size pore, dopant and size of the particles phase [10,11]. For TiO<sub>2</sub> photoinduced super-hydrophilicity, the main efforts have been made in two aspects: one is to narrow the wide bandgap to extend the spectral response of TiO<sub>2</sub> to the visible region for the efficient utilization of the energy from the sun. Another is to reduce the recombination rate of photogenerated electron-hole pairs to enhance efficiency of photolysis. Many efforts have been made to achieve the utilization of

visible light for TiO<sub>2</sub> material, such as transitional metal ion doping [12, 13, 14], non-metal element doping [15, 16] and dye sensitization [17].

In the present study, a series various %wt Cr-doped TiO<sub>2</sub> and undoped TiO<sub>2</sub> have been successfully synthesized using a hot-injection reflux technique. The major goal were to synthesize and characterize undoped TiO<sub>2</sub> and a series various %wt chromium-doped TiO<sub>2</sub> and to investigate its photoinduced super-hydrophilic properties for antifogging materials.

#### 2. MATERIALS AND METHODS

#### 2.1 Materials

Ammonium hydroxide (NH<sub>4</sub>OH, 28-30% NH<sub>3</sub>) solution, hydrogen peroxide solution (H<sub>2</sub>O<sub>2</sub>, 10 wt% in H<sub>2</sub>O), ammonium chromate (NH<sub>4</sub>)<sub>2</sub>CrO<sub>4</sub>, 99%), titanium (IV) chloride (TiCl<sub>4</sub>, 99%) were purchased from Sigma-Aldrich. All the reagents were used without further purification. Titanium dioxide hydrate was obtained from the reaction of TiCl<sub>4</sub> and H<sub>2</sub>O<sub>2</sub> [18]. In a particular procedure, 15 ml TiCl<sub>4</sub> was added into a 500 ml glass flask loaded in an icewaterbath, then 30 ml of H<sub>2</sub>O<sub>2</sub> was added slowly into the reaction vessel under magnetic stirring. The precipitate was filtered, washed with distilled water and dried at 100 °C for 5 hours.

#### 2.2 Sample Preparation

A series of chromium doped TiO<sub>2</sub> at various %wt Cr were prepared by a reflux technique. In a particular procedure, 10 g of titanium dioxide hydrate was dissolved in 50 ml of distilled water under vigorous stirring and was stirred for 4 hours to obtain colloid labeled P. For studying the effect of the (NH<sub>4</sub>)<sub>2</sub>CrO<sub>4</sub> concentration, in a separated beaker 0, 3, 6 and 9 wt% Cr-doped TiO<sub>2</sub> respectively were adopted. It was

dissolved in 20 mL of distilled water thoroughly under vigorous stirring to obtain solutions labeled Q<sub>1</sub>, Q<sub>2</sub>, Q<sub>3</sub>, and Q<sub>4</sub> respectively. Each solution Q<sub>1</sub>, Q<sub>2</sub>, Q<sub>3</sub>, and Q<sub>4</sub> was then slowly added to each solution P. The solution mixture was heated at 150 °C with a magnetic stirrer in equipment reflux, added dropwise NH<sub>4</sub>OH until pH to about 8-10 within about 10 minutes. The solution mixture was refluxed at 150 °C for 6 hours. The precipitate was filtered, washed with distilled water and dried at 70 °C for 3 hours. Furthermore, the precipitate was calcined at 600 °C for 2 hours.

#### 2.3 Physical Measurements of Samples

The morphologies of the prepared materials were observed by a scanning electron microscope (Phenom ProX Desktop SEM) equipped energy dispersive X-ray spectroscopy (EDS). Sample surface was observed and the images were recorded. EDS was used to analyze the presence of Ti, and O elements in the TiO<sub>2</sub> and the presence of Ti, Cr, and O elements in the Cr-doped TiO<sub>3</sub>.

The powder XRD patterns of prepared materials were collected using a Rigaku Miniflex 600-Benchtop X-ray diffractometer, operating in the Bragg configuration using Cu K $\alpha$  radiation ( $\lambda = 1.5406$  Å) at a tube current of 15 mA and a voltage of 40 kV. Data were collected over 2q values from 2 - 90°. The measurements were recorded in steps of 0.02° with a count time of 5 s/step at room temperature 25 °C. The qualitative analysis was carried out with the identification of a phase or phases in the samples by comparison with "standard" patterns: COD and ICDD. The average crystallite size of anatase and rutile were calculated based on XRD peak broadening using the basic Scherrer formula (Eq. (1)) [19], which is then modified and written as Eq. (2). It is modified

by making logarithm on both sides:

$$\beta = \frac{K\lambda}{L\cos\theta} = \frac{K\lambda}{L} - \frac{1}{\cos\theta} \qquad \dots (1)$$

$$\ln \beta = \ln \frac{K\lambda}{L\cos\theta} = \ln \frac{K\lambda}{L} + \ln \frac{1}{\cos\theta}$$
 .... (2)

where L is the average crystallite size,  $\beta$  is the peak width of the diffraction peak profile at half maximum height (FWHM) resulting from small crystallite size in radians and *K* is a constant related to crystallite shape, normally taken as 0.9, l is the wavelength of the X-ray radiation ( $\lambda K_{\alpha}(Cu) = 1.5406 \text{ nm}$ ) and  $\theta$  is the Bragg angle. If we plot the results of  $\ln \beta$  against  $\ln (1/\cos q\theta)$ , then a straight line with a slope of around one and then an intercept of about  $\ln K/L$  must be obtained. The mean crystallite size of anatase particle was estimated by analysing the broadening of the (101), (004), (200), (211), and (220), reflections. While the mean crystallite size of rutile particle was estimated by analysing the broadening of the (110), (101), (111), (210), and (220), reflections. The average crystallite size of brookite and srilankite (TiO2-II) were calculated based on XRD peak broadening using the Scherrer Formula (Eq.(1)).

Full adsorption-desorption isotherms data of nitrogen at 77 K on all prepared materials were collected at various partical pressures in a Surface Area and Pore Porosimetry Analyzer ASAP 2020 instrument from Micromeritics. Before the BET/BJH measurements, the prepared materials were degassed at 150 °C under vacuum for 4 h prior to analysis with a vacuum set point of 10 mmHg. The Brunauer-Emmett-Teller (BET) specific surface data area ( $S_{BET}$ ) was determined by a multipoint BET method using the adsorption data in the relative pressure (P/P0) of  $\sim 0.30$  [20]. The pore size distribution was evaluated from the adsorption-desorption branch of the

isotherms by the procedure developed by Barrett, Joyner and Halenda (BJH) [21]. The nitrogen adsorption and desorption volume at the relative pressure (P/P0) of  $\sim 0.99$  was used to determine the pore volume and the average pore size.

## 2.4 Study of Photoinduced Superhydrophilic Properties

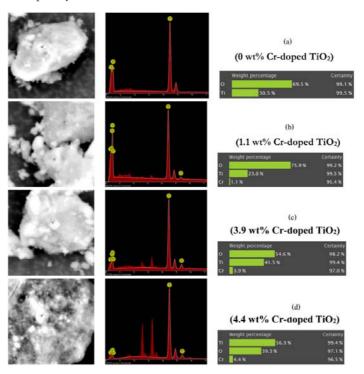
Ultraviolet-ray irradiated to the surface of the prepared sample by commercial 20W

black light blue fluorescent light, and the contact angle of water was measured every 2 minute.

#### 3. RESULTS AND DISCUSSIONS

## 3.1 Scanning Electron Microscopy (SEM)

Figure 1 show typical SEM images and EDS analysis of  ${\rm TiO}_2$  and  ${\rm Cr\text{-}doped\ TiO}_2$  nanoparticles.



**Figure 1.** SEM image (left), EDS analysis (middle) and weight percentage of Ti, O and Cr in the prepared samples (right): (a) undoped TiO<sub>2</sub>, (b) 1.1 wt% Cr-doped TiO<sub>2</sub>, (c) 3.9 wt% Cr-doped TiO<sub>2</sub>, and (d) 4.4 wt% Cr-doped TiO<sub>2</sub>.

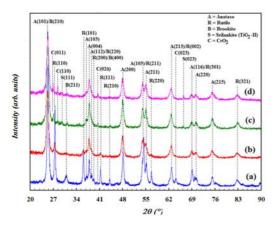
SEM micrographs and EDS spectra of TiO<sub>2</sub> (Figure 1(a)) and 1.1., 3.9, and 4.4 wt% Cr-doped TiO<sub>2</sub> (Figure 1(b-d)) prepared by reflux technique show the formation of aggregated secondary particles by the agglomeration of primary particles. On the theoretical basis, addition of each: 3, 6 and 9 wt% Cr-doped TiO<sub>2</sub> should produce experimentally only 1.1, 3.9 and 4.4 wt%

Cr-doped TiO<sub>2</sub>, respectively. The EDS analysis reveals the presence of Ti and O elements in TiO<sub>2</sub> and the presence of Ti, Cr and O elements in various wt% Cr-doped TiO<sub>3</sub>.

#### 3.2 X-ray Diffraction (XRD)

Figure 2 represents the XRD patterns of undoped TiO<sub>2</sub> and Cr-doped TiO<sub>2</sub>.

The undoped TiO<sub>2</sub> shows that anatase (major), rutile (minor) and brookite (minor) forms are obtained by reflux technique.



**Figure 2.** PXRD pattern of the prepared samples: (a). undoped TiO<sub>2</sub>, (b). 1.1 wt% Crdoped TiO<sub>2</sub>, (c). 3.9 wt% Cr-doped TiO<sub>2</sub>, and (d). 4.4 wt% Cr-doped TiO<sub>3</sub>.

From the XRD pattern (Figure 2(a)), the peak position at  $2q = 25.36 \square 37.84 \square 48.11 \square$ 54.38□, 55.07□, and 62.88□ are indexed as the (101), (103), (200), (105), (211), and (213) reflections of crystalline anatase phase, corresponding to those shown in the ICDD card No. 00-021-1272. The other diffraction peaks are observed  $2q = 27.53 \square$ ,  $36.14 \square$ 39.24 41.32 and 54.38 are indexed as the (110), (101), (200), (111), and (211) reflections of crystalline rutile phase, corresponding to those shown in the COD card No. 9004141. The three distinct diffraction peaks are clearly observed at  $2q = 25.36 \square$ ,  $30.95 \square$ , and  $39.24 \square$ being assigned to (210), (211), and (400) reflections of brookite phase, respectively, corresponding to those shown in the ICDD card No. 00-016-0617. In the 1.1.wt% Cr-doped TiO2, its phase composition are anatase (major) and rutile (minor). The chromium oxide (CrO<sub>2</sub>), anatase (major), brookite and srilankite (TiO<sub>2</sub>-II) are present in the 3.9 wt% Cr-doped TiO2 and the

4.4 wt% Cr-doped TiO<sub>2</sub>.

The phase composition, the average crystallite sizes (L) of the phases in undoped  ${\rm TiO}_2$  and  ${\rm Cr\text{-}doped\ TiO}_2$  are given in Table 1. It is clear that the crystallite size of anatase decreases (114.09-110.17 nm) with increasing the molar of doping agent ( ${\rm Cr}$ ). The crystallite size of anatase and rutile increase with the presence of doping agent ( ${\rm Cr}$ ).

#### 3.3 N<sub>2</sub> Adsorption-Desorption Isotherm

To investigate the pore size distribution and adsorption properties of undoped TiO, and various wt% Cr-doped TiO2, N2 adsorption-desorption isothermal tests were carried out using BET-BJH method, and their isotherm curves were presented in Figure 3. In all prepared materials, it can be observed that the powder exhibits the classical shape of type-IV isotherm curves with H2-type hysteresis loop according to the IUPAC classication [22, 23]. Their narrow hysteresis loops exhibit a typical pattern of Type IV at a relative pressure from 0.68 to 0.98 (undoped TiO2), 0.60 to 0.92 (1.1 wt% Crdoped TiO<sub>2</sub>), 0.42 to 0.92 (3.3 wt% Cr-doped TiO<sub>2</sub>) and 0.45 to 0.90 (4.4 wt% Cr-doped TiO<sub>2</sub>), indicating that the prepared materials have characteristic of a material that contains mesoporosity and has a high energy of adsorption. In addition, the hysteresis loops for these materials are H2 which means that the material is often associated pores with narrow and wide sections and possible interconnecting channels.

The pore size distribution of undoped TiO<sub>2</sub> and various wt% Cr-doped TiO<sub>2</sub> depicted in Figure 4 (inset) show a porosity in the range of 4.95-12.16 nm. The surface area, volume and pore size distribution of the prepared materials (Cr-doped TiO<sub>2</sub> and undoped TiO<sub>2</sub>) have been summarized in Table 2.

**Table 1.** Phase and crystallite size of undoped  ${\rm TiO_2}$  and  ${\rm Cr\text{-}doped\ TiO_2}$ .

		_	2		- 2	
Sample	Phase*)	Hkl	2θ (°)	d (Å)	FWHM 2θ (deg)	L **) (nm)
Undoped TiO <sub>2</sub>		(101)	25.36	3.509	0.42	
		(004)	37.84	2.376	0.43	
	Anatase	(200)	48.11	1.889	0.47	193.84
		(211)	55.07	1.666	0.47	
		(220)	70.34	1.337	0.50	
		(110)	27.53	3.237	0.25	
		(101)	36.14	2.483	0.29	
	Rutile	(111)	41.32	2.183	0.24	323.83
		(210)	44.13	2.050	0.26	
		(220)	56.63	1.624	0.29	
	Brookite	(211)	30.95	2.887	0.67	122.99
		(101)	25.29	3.519	0.72	
		(004)	37.75	2.381	0.74	
	Anatase	(200)	48.00	1.894	0.70	114.09
		(211)	55.12	1.665	0.49	
1.1 wt% Cr-doped TiO <sub>2</sub>		(220)	70.18	1.340	0.86	
		(110)	27.45	3.247	0.47	
	Rutile	(101)	36.05	2.489	0.52	178.04
		(111)	41.24	2.187	0.51	
		(101)	25.31	3.516	0.73	
		(004)	37.92	2.371	0.76	
3.9 wt% Cr-doped $\mathrm{TiO}_2$	Anatase	(200)	48.01	1.894	0.77	110.17
		(211)	55.03	1.667	0.79	
		(220)	70.15	1.341	0.84	
	Brookite	(211)	31.13	2.870	0.28	294.42
		(011)	27.11	3.287	0.18	
	CrO,	(110)	28.40	3.139	0.10	572.86
	2	(020)	40.56	2.222	0.19	
	TiO <sub>2</sub> -II	(111)	29.62	3.013	0.44	186.69
4.4 wt% Cr-doped TiO <sub>2</sub>		(101)	25.33	3.513	0.72	
		(004)	37.83	2.377	0.74	
	Anatase	(200)	48.01	1.894	0.76	112.59
		(211)	55.06	1.667	0.80	
		(220)	70.11	1.341	0.84	
	Brookite	(211)	31.29	2.8567	0.12	687.24
		(011)	27.06	3.292	0.11	
	$CrO_2$	(110)	28.39	3.141	0.20	506.69
	2	(020)	40.43	2.229	0.23	
	TiO,-II	(111)	29.53	3.022	0.39	210.58
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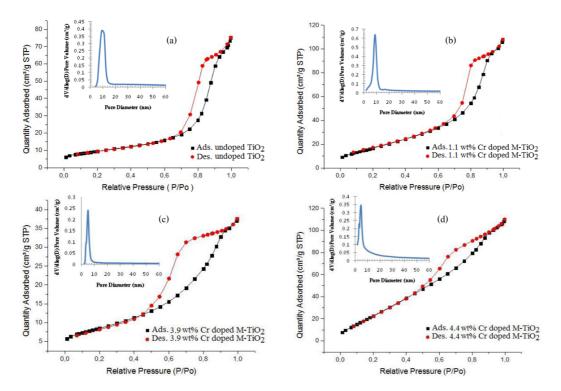
<sup>\*)</sup> The phase composition was determined by qualitative analysis ("standard" patterns: COD and ICDD)
\*\*) The average crystallite size of anatse and rutile were calculated by Modified Debye-Scherrer formula, while for CrO<sub>2</sub>, brookite and srilankite were calculated by Debye-Scherrer formula

Table 2. Surface area, volume and pore size distribution of Cr-doped $\Pi O_2$ and undoped
${ m TiO_2}$ from Nitrogen Adsorption-desorption Isotherm Measurements.

Sample	Surface Area BET	Pore Volume at P/P <sub>o</sub>	Pore Size
	$(S_{\rm BET}) \ ({\rm m}^2/{\rm g})$	$\approx 0.99 \text{ (cm}^3/\text{g)}$	(nm)
Undoped TiO <sub>2</sub>	33	0.1168	12.16
1.1 wt% Cr-doped TiO <sub>2</sub>	65	0.1658	8.01
3.9 wt% Cr-doped TiO <sub>2</sub>	30	0.0591	6.88
4.4 wt% Cr-doped TiO <sub>2</sub>	111	0.1612	4.95

The BET surface area and the mean pore size of the 4.4 wt% Cr-doped TiO<sub>2</sub> exhibit a maximum surface area of 111 m<sup>2</sup>/g, corresponding to mean porous size of 4.95 nm. The pore size distribution curve calculated from the desorption branch of

the isotherm BJH analyses shows that the undoped TiO<sub>2</sub> exhibits pore size of 12.16 nm and the 1.1, 3.9, and 4.4 wt% Cr-doped TiO<sub>2</sub>) exhibit pore sizes of 8.01, 6.88, and 4.95 nm (inset Figure 3), respectively.



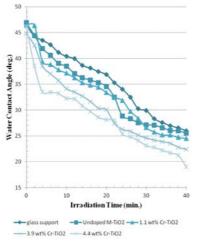
**Figure 3.** Nitrogen adsorption-desorption isotherms of the prepared samples: (a) undoped TiO<sub>2</sub>, (b) 1.1 wt% Cr-doped TiO<sub>2</sub>, (c) 3.9 wt% Cr-doped TiO<sub>2</sub>, and (d) 4.4 wt% Cr-doped TiO<sub>2</sub> (inset of pore size distribution from the adsorption branch of isotherm).

# 3.4 The Contact Angle Changes of Water on the Cr-doped TiO<sub>2</sub> Surface Irradiated by Ultra-Violet (UV) Light

Figure 4 shows diagram of the change of contact angles of water dropped on the UV-iradiated undoped TiO<sub>2</sub> and Cr doped TiO<sub>2</sub> films. The contact angle of water on the prepared samples surfaces can be altered by UV irradiation.

From the diagram in Figure 4, it can be seen that the 4.4 wt% Cr-doped TiO<sub>2</sub> has the most excellent superhydrophilic properties as compared with other samples. The phenomena after UV irradiation for 40 minutes are as follows:

- -Undoped TiO<sub>2</sub>, irradiation with UV light inducing a decrease in contact angle from about 46.93° to 25.51°.
- 1.1 wt% Cr-doped TiO<sub>2</sub>, irradiation with UV light inducing a decrease in contact angle from about 46.48° to 24.52°.
- 3.9 wt% Cr-doped TiO<sub>2</sub>, irradiation with UV light inducing a decrease in contact angle from about 44.84° to 22.38°.
- 4.4 wt% Cr-doped TiO<sub>2</sub>, irradiation with UV light inducing a decrease in contact angle from about 46.12° to 19.04°.



**Figure 4.** Change of the contact angles for film of glass support, undoped TiO<sub>2</sub>, 1.1 wt% Cr-doped TiO<sub>2</sub>, 3.9 wt% Cr-doped TiO<sub>2</sub>, and 4.4 wt% Cr-doped TiO<sub>2</sub> with UV irradiation.

#### 4. CONCLUSIONS

A various wt% chromium-doped TiO, (Cr-doped TiO<sub>2</sub>) and undoped TiO<sub>2</sub> have been successfully synthesized by hot-injection reflux technique. The prepared samples consist of anatase (major), rutile (minor), chromium oxide (CrO<sub>2</sub>) (minor), brookite (minor) and srilankite (TiO<sub>2</sub>-II) (minor) type structures. In the 1.1.wt% Cr-doped TiO<sub>2</sub>, its phase composition is anatase (major) and rutile (minor). The CrO<sub>2</sub>, anatase, brookite and srilankite (TiO<sub>2</sub>-II) are present in the 3.9 wt% Cr-doped TiO2 and the 4.4 wt% Cr-doped TiO<sub>2</sub>. The BET surface area and the mean pore size of the 4.4 wt% Cr-doped TiO, exhibit a maximum surface area of 111 m<sup>2</sup>/g, corresponding to mean porous size of 4.95 nm. All prepared materials (Cr-doped TiO<sub>2</sub> and undoped TiO<sub>2</sub>) exhibit mesoporous of type-IV isotherm curves with H2-type hysteresis loop according to the IUPAC classication. The hydrophilic properties of Cr-doped TiO, were investigated with illumination of UV light, and all prepared samples show excellent super-hydrophilic properties, and the 4.4 wt% Cr-doped TiO, exhibits the most excellent superhydrophilic properties as compared with other samples. These results allow the materials to be prospective application as antifogging.

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