

Submicron-Lubricant Based on Crystallized Fe_3O_4 Spheres for Enhanced Tribology Performance

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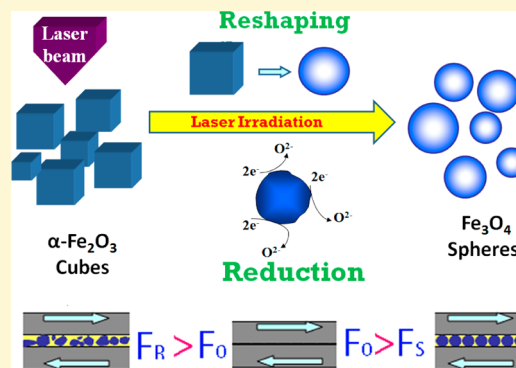
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S Supporting Information

ABSTRACT: We demonstrate in this paper a novel finding of submicron-lubricant exemplified by crystallized magnetite (Fe_3O_4) submicrometer spheres. Optimized tribological performance of lubricant particles occurs when the particles are submicrometer sized and spherical. The tribology of lubricant particles is thus verified to be size and shape dependent. Owing to the unique heating and quenching process involved in pulsed laser irradiation, Fe_3O_4 submicrometer spheres are acquired through phase transformation and reshaping. The formation process of the regular Fe_3O_4 spheres in deionized water was investigated by choosing different laser fluences and irradiation times. The shape and phase transformation mechanisms are elucidated as well. Tribological properties of the obtained Fe_3O_4 submicrometer spheres have been explored, inspired by their spherical shape and smooth surfaces. Crystallized magnetite submicrometer spheres under optimized conditions show a reduction in friction and wear by up to 40% and 20%, respectively. Control experiments revealed that nanoparticles, nonspherical submicrometer particles, and micrometer particles could not reach such good tribology performance, indicating the potential of adopting crystallized Fe_3O_4 submicrometer spheres in the application of mechanical energy saving.



1. INTRODUCTION

The ability to generate micro/nanocrystals with well-controlled sizes, shapes, and compositions is central to advance applications ranging from catalysis to electronics, photonics, sensing, as well as energy generation, storage, and conversion.^{1–5} Developing nanostructures with designed morphologies for investigation of tribology has recently been of importance as one-third of the total mechanical energy used in this world goes waste to counteract the friction force.^{6–8} Nanoparticles including ZnO ^{6,8} and Al_2O_3 ⁷ have shown good potential as additives for lubricant to improve its tribology property. Intriguingly, earth abundant Fe_2O_3 and Fe_3O_4 are more attractive for such applications when considering their high hardness (1030 and ~ 450 Hv, respectively). Attempts have been thus reported by using iron oxides as additives in base oil in the form of micro/nanoparticles.⁹ Numerous methods have been developed for the synthesis of high-quality iron oxide nanoparticles, such as precursor-templated methods,¹⁰ hydrothermal methods,¹¹ and chemical precipitation.¹² However, the morphologies of nanoparticles prepared by the above methods are usually nonspherical.^{13–15} In addition, research related to the particles based lubricant has always

focused on seeking ideal material species,^{6–15} while few on the investigation of size-dependent tribology properties, which is of importance not only fundamentally but for practical application.

In view of the primary principles of lubrication, spherical particles with high hardness are highly appreciated to change sliding into rolling friction more effectively and reduce the wear more significantly. No efforts, however, to the best of our knowledge, have been reported on spherical iron oxides particles for lubrication. As a matter of fact, synthesis of iron oxide particles with spherical shape has been well documented in literatures, however, the acquired particles are dominated by hierarchical structures; that is, the spherical particles are constructed by secondary subunits.¹³ As aforementioned, these particles also are not ideal for lubricating when considering their surface roughness and low hardness. Unfortunately, synthesis of inorganic highly crystalline spheres with smooth surfaces has rarely been reported, possibly because

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of their very high melting points and anisotropic crystal structures.

Laser ablation has recently been adopted to synthesize particles both in vapor or liquid medium, which is known as pulsed laser ablation (PLA). The combination of laser ablation and vapor transport deposition provide good control over the temperature, pressure and gas to grow nanostructures, especially for one-dimensional nanowires or nanotubes.^{16–19} Laser ablation in liquid media was first implemented to fabricate colloidal solutions of nanoparticles, known as laser ablation in liquid (LAL).^{20,21} Henglein et al.²² first applied a pulsed laser to ablate pure metal targets in various solvents to grow colloidal solutions containing metal nanoparticles (NPs). Since then, LAL has been deeply developed as an important method to prepare metal, semiconductor, and even polymer NPs.^{23–25} For instance, Zeng et al.^{26–28} reported composition-controlled synthesis of ZnO-Zn composite nanoparticles by laser ablation of a zinc metal target in pure water and aqueous solution of sodium dodecyl sulfate and studied their optical properties; Yang et al.^{29,30} have synthesized Si quantum dots and SiC quantum rings using LAL and an anomalous quantum confinement effect was observed; Jayawardena et al.³¹ showed that ZnO nanocrystals and nanorods can be fabricated by hybrid hydrothermal and UV laser heating growth process.

Most recently, metal and semiconductor submicrometer spheres have been prepared by a selective pulsed laser heating process with proper laser wavelength and energy,^{32,33} which effectively inhibits the anisotropic crystal growth and favors for acquiring inorganic crystalline spheres. Inspired by this, in this manuscript, we introduce crystalline Fe₃O₄ submicrometer spheres for lubrication by simple pulsed laser irradiation of preformed α -Fe₂O₃ cubic particles. A laser-stimulated surface tension energy release (LSTER) and chemical phase transition process involved in the laser irradiation process assures the formation of spherical Fe₃O₄ particles. What is more important, the lubrication of iron oxide NPs, cubic submicrometer particles and microparticles was further investigated, verifying a novel phenomenon of “submicron-lubricant”, that is, crystalline submicrometer spheres exhibiting superior tribology performance.

2. EXPERIMENTAL SECTION

2.1. Synthesis of Hematite (Fe₂O₃) Cubic Particles. Fe₂O₃ cubic particles were prepared by a hydrothermal process. In a typical reaction, 0.729 g CTAB, 0.622 g FeCl₃·6H₂O, and 0.4 g PEG-6000 was dissolved into 60 mL deionized water. The whole mixture was stirred to form a homogeneous solution and then transferred into a Teflon-lined stainless steel autoclave. Then, it was sealed and maintained at 120 °C for 12 h. After the reaction, the precipitated powders were collected by centrifuging and washed several times with deionized water and ethanol. Finally, the samples were dried at 50 °C for 12 h.

2.2. Synthesis of Magnetite Submicrometer Spheres. The as-prepared Fe₂O₃ cubic particles were dispersed into deionized water at a mass concentration of 0.5 wt % to form a colloidal solution. A KrF excimer laser (10 Hz, 25 ns, Coherent, CompexPro 205) was used as the light source for pulsed laser irradiation. The pulsed laser beam was guided by a reflection mirror and focused by a quartz lens on colloidal solution. The solution was then irradiated with a focused laser beam with different laser fluences (400, 700, 1000, 1300 mJ pulse⁻¹ cm⁻²), which were determined by a laser FieldMaxII Meter (Coherent). Laser irradiation of raw Fe₂O₃ particles was performed for different times. After laser irradiation, all prepared colloidal suspensions were centrifuged at 8000 rpm and finally dried at 50 °C for further characterization.

2.3. Structure and Tribology Characterizations. The sample morphology was characterized with a scanning electron microscope (SEM, FEI Quanta 250 FEG). The phase and crystal structures of the particles were investigated with X-ray diffraction (XRD, Cu K α radiation, Bruker D8 ADVANCE) and X-ray photoelectron spectroscopy (XPS, ThermoFisher, ESCALAB 250). The tribological properties of the prepared particles were measured in terms of friction coefficients and wear scar diameters with a tribology tester (MMU-10G, Sinomach-Jinan). Fe₃O₄ submicrometer spheres dispersed into oil with different mass concentration of 0, 0.03, 0.07, 0.13, and 0.2 wt %, respectively, were tested. Control experiments were performed by measuring the tribological properties of as-prepared α -Fe₂O₃ cubic particles, magnetite nanoparticles, hierarchical magnetite spherical particles with 0.03–0.2 wt %. Detailed experimental conditions of the four-ball wear test were set as follows: rotation speed at 1450 rpm, load of 147 N, temperature at 75 °C and time of 30 min. The friction coefficient was recorded *in situ* and the test data were acquired automatically with a computer. The size of the wear scar diameter was observed with a metallographic microscope and then the average size of wear scar diameter was obtained.

3. RESULTS AND DISCUSSION

3.1. Morphology and Phase Analysis of Fe₃O₄ Submicrometer Particles.

Figure 1a shows a typical SEM

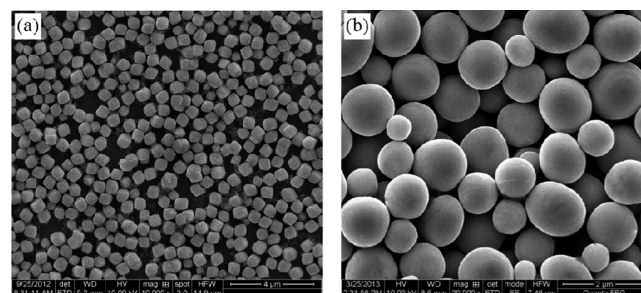


Figure 1. SEM images of (a) nanoparticles prepared by hydrothermal method and (b) submicrometer spheres after irradiation with a 248 nm KrF excimer laser for 60 min under fluence of 1300 mJ pulse⁻¹ cm⁻².

image of as-prepared particles synthesized by a hydrothermal method, from which it can be seen that the particles are cubic with edge length of about 600 nm. After UV laser irradiation for 55 min with fluence of 1300 mJ pulse⁻¹ cm⁻² in deionized water, these cubes transform to submicrometer spheres as shown in Figure 1b. One can note that the size of the spherical particles after laser irradiation is similar to or a little bigger than that of the corresponding raw cubic particles. This is totally different from the case of traditional LAL that a solid or pressed target is laser ablated in liquid, which usually produces much smaller particles with size of a few nanometers.¹⁰ The composition and phase purity of the particles with different morphologies were further examined by XRD, as shown in Figure 2a. The XRD pattern of the as-prepared cubic samples can be well ascribed to the rhombohedral α -Fe₂O₃ (PDF: 24-0072). However, after laser irradiation, the XRD pattern of the submicrometer spheres exhibits different diffraction characteristics. By comparing the XRD peaks with the standard powder diffraction cards, the submicrometer spheres can be identified as the γ -Fe₂O₃ (PDF: 24-0081)³⁴ or Fe₃O₄ (PDF: 89-4319).

XPS measurement was employed to check the phase change upon laser irradiation due to its sensitivity to surface chemical changes. Figure 2b shows the wide scanning spectrum and the photoelectron peaks at binding energies of about 710.8, 530.8,

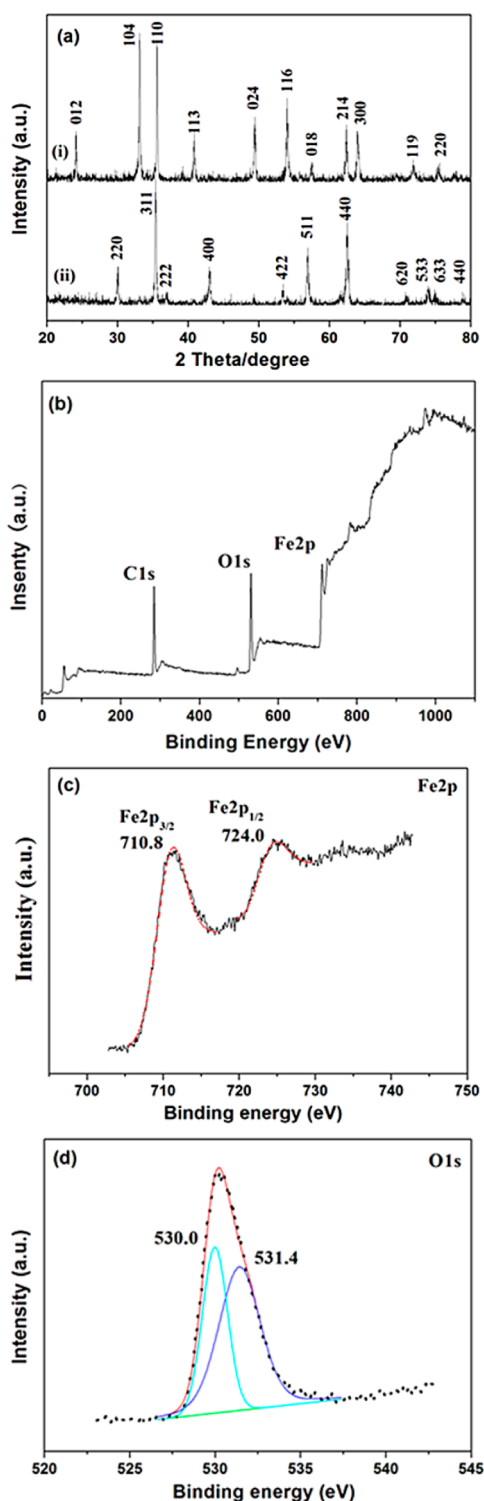


Figure 2. (a) XRD spectra of (i) as-prepared cubic particles by the hydrothermal method, (ii) submicrometer spheres after irradiation with a 248 nm KrF excimer laser for 60 min. (b–d) XPS spectra of submicrometer spheres (b) wide scan, (c) Fe_{2p} spectra, and (d) O_{1s} spectra.

and 285.0 eV, which is attributed to Fe 2p, O 1s, and C 1s, respectively.¹⁵ Figure 2c shows the XPS signals of the Fe 2p peak, with two peaks of Fe 2p_{1/2} at 710.8 eV and Fe 2p_{3/2} at 724.0 eV with the full width at half-maximum (fwhm) of 4.1 and 5.1 eV, respectively. The absence of the satellite peak situated at about 719 eV, which is a major characteristics of Fe³⁺

in γ -Fe₂O₃, clearly excludes the formation of γ -Fe₂O₃ after laser irradiation.³⁵ The O core level spectrum together with their spectral deconvolution curves in Figure 2d consists of a main peak originating from the oxygen in Fe₃O₄ (at 530.0 eV) and a shoulder centered at 531.4 eV, which is ascribed to surface traps.³⁶ Moreover, the total Fe/O ratio was estimated to be 0.74 by curve resolution analysis, which is well consistent with the stoichiometric ratio of Fe₃O₄. The XPS and XRD analysis reveal that the raw cubic particles are rhombohedral α -Fe₂O₃, and spinel Fe₃O₄ submicrometer spheres are obtained after pulsed laser irradiation. Therefore, the physical reshaping and phase transition from α -Fe₂O₃ cubes to Fe₃O₄ spheres are induced by a simple laser irradiation process in water. Compared with the conventional chemical routes, this method is very fast, moderate, and safe, and does not use any surfactants.^{26,32} To disclose more details about the morphology evolution, irradiation time-dependent and laser energy-dependent experiments were further conducted.

3.2. Effect of Laser Fluence and Irradiation Time on Particle Shape.

Figure 3 shows SEM images of the prepared

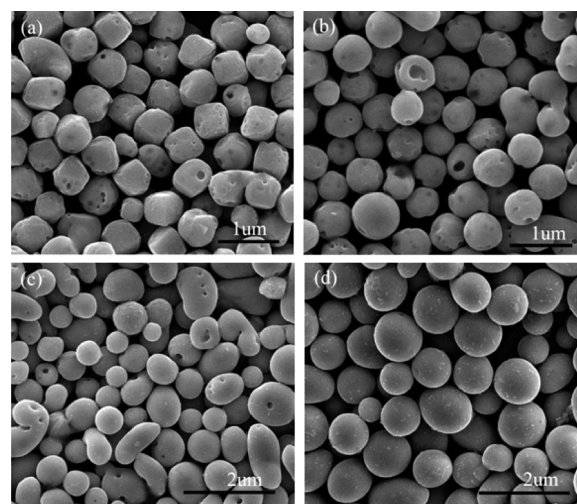


Figure 3. SEM images of the product after irradiating Fe₂O₃ cubic particles by a laser with different fluences for 55 min, (a) 400 mJ pulse⁻¹ cm⁻², (b) 700 mJ pulse⁻¹ cm⁻², (c) 1000 mJ pulse⁻¹ cm⁻², and (d) 1300 mJ pulse⁻¹ cm⁻².

Fe₃O₄ particles. When the laser fluence is as low as 0.4 J pulse⁻¹ cm⁻², the particles do not show clear morphological changes, but only few small spherical particles are formed, as shown in Figure 3a. This is because the laser cannot produce enough energy to melt the particle surfaces. When the laser fluence is increased to 0.7 J pulse⁻¹ cm⁻², all particles exhibit morphologies different from those of the original ones. A number of small particles transform to spheres (Figure 3b). When the laser fluence is further increased, the melting of the particle surfaces and then merging of different particles are clearly exhibited, as shown in Figure 3c. α -Fe₂O₃ particles change gradually from cube morphology to regular spherical particles when the laser energy density is further increased to 1300 mJ pulse⁻¹ cm⁻² (Figure 3d). This indicates that lasers with proper higher fluence are necessary for submicrometer sphere formation. Moreover, during the particle shape transformation and coalescence processes, the volume of the particle(s) remains constant but the surface areas decrease in comparison with the

original particle(s). Therefore, the high surface energy (E_s) of particles is released as EST is proportional to the surface area.

Figure 4 further reveals the morphology evolution of α - Fe_2O_3 particles with irradiation time for the same laser fluence

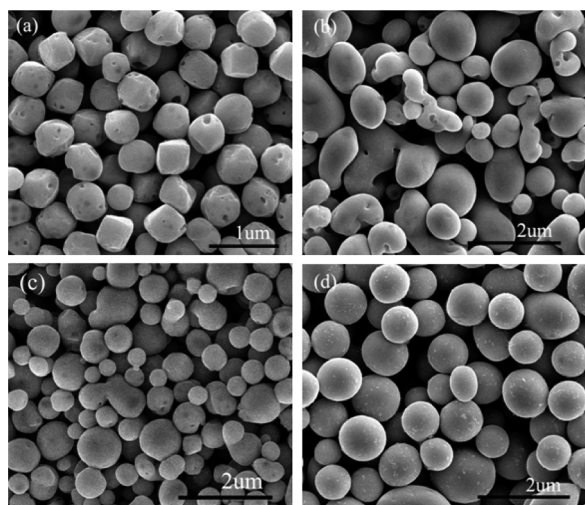


Figure 4. SEM images of the product after irradiating Fe_2O_3 cubic particles with a laser fluence of $1300 \text{ mJ pulse}^{-1} \text{ cm}^{-2}$ for different times: (a) 10 min; (b) 25 min; (c) 40 min; and (d) 55 min.

($1.3 \text{ J pulse}^{-1} \text{ cm}^{-2}$). As the applied laser energy density is high enough to melt the particles, some cubic particles begin to lose their straight edges after 10 min laser irradiation, as shown in Figure 4a. Quasi-spherical particles emerge after 25 min irradiation, as shown in Figure 4b. Meanwhile, particle merging is also observed; that is, several particles begin conjugating into bigger ones. While further prolonging laser irradiation, spherical Fe_3O_4 particles are formed, as shown in Figure 4c and d. Both particle rounding and coalescence reduce their surface area and, accordingly, the release of the surface energy.

3.3. Mechanisms of Morphology Evolution and Phase Transformation. The results above indicate that the reshaping and phase transformation of particles can be delicately controlled by laser fluence and irradiation time. Both are conducted during pulsed laser irradiation, which experiences typical extreme nonequilibrium conditions in nature such as high temperature (10^4 K), high pressure (1 GPa), and high kinetic energy (1 eV).³⁷ The particle morphology change is due to the high temperature of the particle surface caused by laser irradiation. The key feature of micro/nanoparticles is the high surface area per mass (of order $50\text{--}300 \text{ m}^2/\text{g}$) and one of the consequences of this intrinsic characteristics is the high surface energy concentrated in the surface layers of the particles.²³ The surface energy is proportional to the surface area and can be released by minimizing the surface area. Therefore, the spherical structure has the smallest surface area among all surfaces enclosing a given volume and, accordingly, smallest E_s for spherical particles. Therefore, the pulsed laser irradiation induced heating causes a thermal instability (melting) of the α - Fe_2O_3 cube particle surfaces, cube particles change into spheres gradually with E_s release, as schematically shown in Figure 5. At the same time, since a laser with pulse duration of 25 ns was used, rapid quenching happened in water after pulsed heating. This is helpful to inhibit the reorientation of surface atoms on a spherical particle, and finally, the particles retained the overall spherical morphology. Moreover, if the surface atoms are

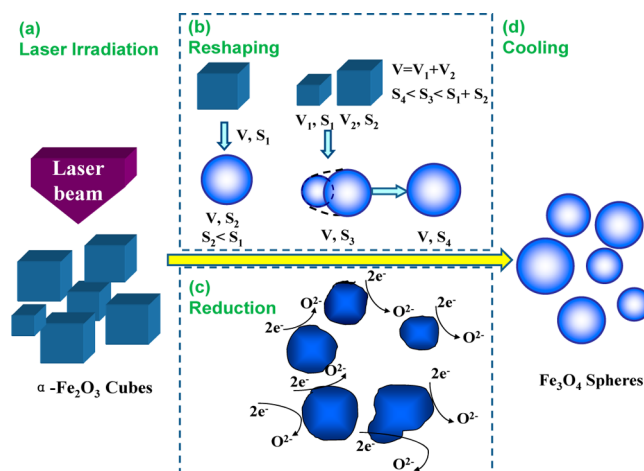
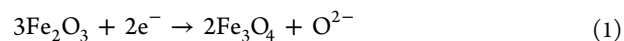


Figure 5. Schematic demonstration of the pulsed laser-induced reshaping (b) and reduction (c) mechanism.

activated by laser irradiation above the interparticle surface potential barrier, contacting particles begin to merge (Figures 3c, 4b) as the surface area of one unified particle is smaller than that of two original individual particles, as shown in Figure 5b. The particle shape change can thus be well described with the above proposed laser stimulated surface energy release growth mechanism. This method is different from the traditional pulsed laser ablation in liquid, in which particle formation is due to the condensation of laser ablated plasma plumes.⁶

The above results including SEM, XRD, and XPS analysis indicate that the α - Fe_2O_3 particle reshaping was mainly due to the high temperature and high cooling rate involved in pulsed laser irradiation. Meanwhile, electrons can also be generated in water under such extreme nonequilibrium conditions.³⁸ The Fe_2O_3 particles get these electrons and then release oxygen ions simultaneously, as shown in Figure 5c. Some of the Fe^{3+} ions in the lattice were thus reduced to Fe^{2+} ions to maintain electroneutrality. Two oxygen ions may recombine to form an O_2 monomer with electron released. These released electrons bound by other Fe_2O_3 particles finally transformed all Fe_2O_3 particles into Fe_3O_4 . This process can be simply described by the two chemical reactions 1 and 2. Therefore, the transformation between these two iron oxides is thought to be determined by the diffusion of Fe^{2+} and Fe^{3+} ions within the oxygen sublattice and electron transfer between different valences of Fe ions.^{39–41}



3.4. Fe_3O_4 Submicrometer Spheres with Enhanced Tribology Performance. Inspired by their spherical shape and smooth surfaces, the tribology properties of Fe_3O_4 submicrometer spheres (shown in Figure 1b) were measured by adding them in commercial lubricant oils in terms of friction coefficients and diameters of wear scar, while taking cubic submicrometer particles (Figure 1a), nanoparticles (Supporting Information, Figure S1a), and microparticles (Supporting Information, Figure S2a) as control samples. Figure 6 shows the variation of friction coefficients with time when the commercial pristine lubricating oil (Curve marked with 0 wt %) and lubricating oil with α - Fe_2O_3 cube particles (Figure 6a) or Fe_3O_4 submicrometer spheres (Figure 6b) as additives are

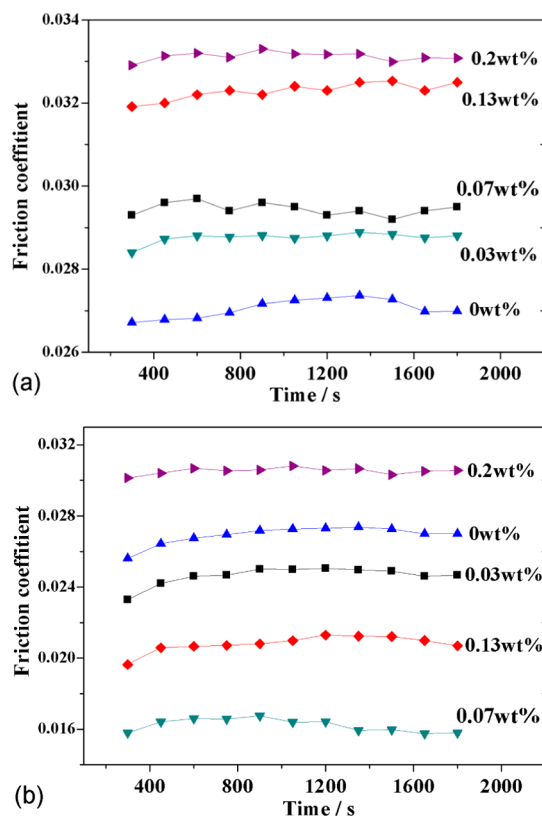


Figure 6. Friction coefficient curve measured with a four-ball tester when particles of different mass concentration were added into the tested oil, respectively, (a) as-prepared Fe₂O₃ cubes and (b) submicrometer spherical Fe₃O₄ particles. The friction coefficient measured with pure commercial lubricating oil (curve marked with 0 wt %) is also shown for comparison.

measured with a four-ball tribology tester. It can be seen that the friction coefficients increase obviously when Fe₂O₃ cubic submicrometer particles (0.03–0.2 wt %) are added into commercial lubricating oil. The reason is that Fe₂O₃ particles with cubic morphology depositing onto the thrust-rings increase the sliding friction. However, when a certain amount of Fe₃O₄ submicrometer spheres are added into lubricating oils, the friction coefficients apparently decrease. The friction-reduction properties show a clear dependence on the concentration of Fe₃O₄ spherical particles, as shown in Figures 6a. The friction coefficients decrease when the concentration is smaller than 0.15 wt %. However, the friction coefficients begin to increase when the Fe₃O₄ concentration is larger than 0.2 wt %. Therefore, when the amounts of submicrometer spheres reach an optimal concentration, obvious friction-reduction effect is acquired.

For Fe₃O₄ spherical particles, when the sphere concentration is 0.07 wt %, the average friction coefficient decreases by about 38%, which is much more significant than other reported reduction data (<20%).⁴²

Antiwear ability is another important characteristic index for lubricant property. Therefore, the wear scar diameter (WSD) was also measured during tribological tests. A metallographic microscope was used to investigate the antiwear property of different lubricating oils. Figure 7 compares two typical wear scar images of the testing balls in lubricant oil and oil with 0.07 wt % Fe₃O₄ submicrometer spheres, showing a clear decrease of the WSD. Table 1 compares the average wear scar diameters on

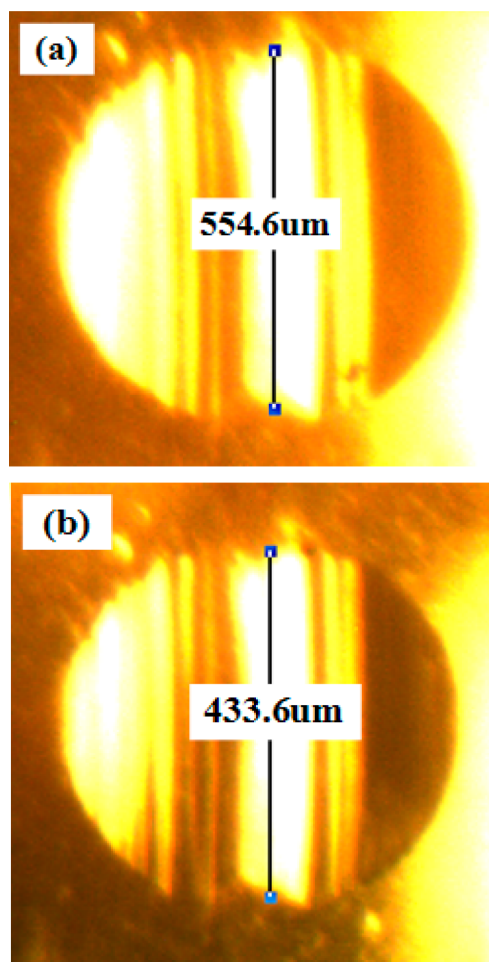


Figure 7. Wear scar images after the four ball test. (a) Lubricated with pure oil and (b) lubricated with oil involving 0.07 wt % Fe₃O₄ submicrometer spheres.

the testing balls lubricated by lubricant oil dispersions of submicrometer cubic particles and spheres with different mass concentrations. Due to the intrinsic high hardness, both of the particles exhibit antiwear properties under optimized concentrations, which is similar to that of Al₂O₃ nanoparticles.^{25,43} The lubricant oil dispersion with 0.07 wt % Fe₃O₄ spheres performs the most effective, that is, with WSD reduced by 22%. Similar to the friction-reduction effect, higher concentration of Fe₃O₄ submicrometer spheres or cubic particles (e.g., 0.2 wt %) will deteriorate the antiwear performance. The maximum decrease of the average friction coefficient and WSD is about 38% and 22%, respectively, when an optimized concentration of Fe₃O₄ submicrometer spheres reaches 0.07 wt %.

Further control experiments reveal that nanoparticles and microparticles could not reach such good tribology performance from submicrometer spheres. The friction coefficients decreased when Fe₃O₄ NPs were added into lubricating oil and the average friction coefficient is smallest when the concentration is 0.1 wt %. However, the average friction coefficient decreases by only about 10% (Supporting Information, Figure S1b), which is much smaller than for the Fe₃O₄ submicrometer spheres (38%) reported in the paper. Moreover, the best WSD reduction (10%) is also smaller than that of submicrometer spheres (22%), as seen in Supporting Information, Table S1. In comparison with the submicrometer Fe₃O₄ spheres, all friction coefficients (Supporting Information,

Table 1. Wear Scar Diameter on the Testing Balls Lubricated with Different Lubricating Oil Involving Fe₂O₃ Cubic (C) Particles or Fe₃O₄ Submicrometer Spheres (S) with Different Mass Concentrations

Fe ₂ O ₃ /Fe ₃ O ₄ concn. (wt %)	0	0.03C	0.07C	0.13C	0.2C	0.03S	0.07S	0.13S	0.2S
avg. with error (μm)	555 $\pm$ 4	535 $\pm$ 5	504 $\pm$ 4	606 $\pm$ 3	623 $\pm$ 3	532 $\pm$ 2	434 $\pm$ 5	538 $\pm$ 2	583 $\pm$ 4
reduction (%)		4	9	−9	−12	4	22	16	−5

Figure S2b) and WSD (see Supporting Information, Table S2) increased when Fe₃O₄ microparticles were added into commercial lubricating oil.

The exact mechanism of nanoparticle action as lubricant for enhancing tribological performance is still not very clear.^{24,43,44} Basically, inorganic particles are expected to serve as spacer and form a lubrication film when dispersing them in lubricant oils.⁴⁰ With the oil is flowing during friction, the nanoparticles can stay between the rubbing surfaces. The existence of nanoparticles on the thrust-ring surface acting as spacer to reduce the friction between the thrust-rings has been identified with SEM and energy-dispersive X-ray spectrum.⁴³ When the shape of the Fe₃O₄ particles is spherical, the friction between thrust-rings will change from sliding friction to rolling friction using the spheres as molecular bearings.^{44–46} As a result, the friction coefficients reduce prominently, as shown in Figure 6b. However, the Fe₂O₃ particles with cube shapes have no such effect and play a negative role on friction reduction, leading to the friction coefficient increase. Therefore, the friction performance of lubricating oil with different Fe₃O₄ submicrometer sphere concentrations can be primarily understood. When the concentration of Fe₃O₄ spheres is low, a small amount of spherical particles starts to play the role of rolling friction instead of sliding friction, which explains well the decrease of the friction coefficients when the additive concentration increases. However, if the concentration of spheres is too high, the redundant spheres will aggregate and chemically condense, which severely enhances friction. Therefore, there is an optimal concentration of submicrometer spheres for lubrication (e.g., 0.07 wt % for Fe₃O₄ spheres in our study).

The enhanced antiwear performance of lubricating oil can be explained with the self-repairing mechanism for lubricant oils containing inorganic particles.^{46,47} Fe₂O₃ or Fe₃O₄ particles deposited onto the worn steel surface under compressive stress during the friction process could form a protective film and act as spacers between two friction counterparts. It prevents the metal-to-metal direct contact and fills the wearing scars. In this case, the high hardness of iron oxide is an intrinsic characteristic of the antiwear application. However, the antiwear ability is also affected by the particle concentration, and when the Fe₃O₄ microsphere additive concentration reached 0.07 wt %, a better antiwear performance of lubricant oil in terms of WSD was obtained.

4. CONCLUSIONS

In summary, a novel submicron-sized lubricant based on crystalline iron oxide spheres is demonstrated in this paper revealing that optimized tribology performance can be acquired when lubricant particles are spherical and submicrometer sized. The magnetite submicrometer spheres have been synthesized by laser irradiation hematite cubic particles in water via edge rounding and/or coalescence together with a phase transition. Upon laser irradiation, the Fe₂O₃ particle surface was thermally activated (melting) and the high surface energy concentrated in the surface layer will be released when cube particles change

into spheres gradually as a result of surface area minimization, during which electrons in water reduce partially Fe³⁺ to Fe²⁺ ions under a transient high temperature process by laser irradiation, leading to a phase transition from α -Fe₂O₃ to Fe₃O₄. Inspired by their high hardness and smooth surface of the magnetite submicrometer spheres, their tribology performance was explored. This showed that the Fe₃O₄ oxide submicrometer spheres, different from Fe₂O₃ cubic particles, can more effectively reduce the friction coefficients and wear scar diameters under an optimized concentration when dispersing them in lubricating oil, which is a typical morphology-dependent tribological application. Control tribology studies on nanoparticles and micrometer particles revealed that the lubricant property can be optimized when the particles sizes are submicrometer scaled. It is also worth noting that the method applying laser irradiation in liquid medium might be conveniently extended to design other oxide spheres, compensating for the shortcomings of traditional chemical methods in preparing these nanomaterials. We thus expect that our study will be of significance to advance studies on submicrometer sphere based mechanical energy saving applications.

■ ASSOCIATED CONTENT

§ Supporting Information

Tribology properties of Fe₃O₄ nanoparticles, Fe₃O₄ microparticles with irregular shapes, and Fe₂O₃ particles and their size distribution after irradiation with different energy densities. This material is available free of charge via the Internet at <http://pubs.acs.org>

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Notes

The authors declare no competing financial interest.

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■ REFERENCES

- (1) Xia, Y. N.; Xia, X. H.; Wang, Y.; Xie, S. F. *MRS Bull.* **2013**, *38*, 335–344.
- (2) Yang, S. K.; Cai, W. P.; Kong, L. C.; Lei, Y. *Adv. Funct. Mater.* **2010**, *20*, 2527–2533.

- (3) Oh, M. H.; Yu, T.; Yu, S. H.; Lim, B.; Ko, K. T.; Willinger, M. G.; Seo, D. H.; Kim, B. H.; Cho, M. G.; Park, J. H.; Kang, K.; Sung, Y. E.; Pinna, N.; Hyeon, T. *Science* **2013**, 340 (6135), 964–968.
- (4) Zhang, L.; Blom, D. A.; Wang, H. *Chem. Mater.* **2011**, 23 (20), 4587–4598.
- (5) Kensuke, A.; Satoshi, A.; Takaaki, T.; Shingo, I.; Satoshi, T.; Hidemi, N. *Chem. Mater.* **2008**, 20 (9), 3042–3047.
- (6) Hernández Battez, A.; González, R.; Viesca, J. L.; Fernández, J. E.; Fernández, J. M. D.; Machado, A.; Chou, R.; Riba, J. *Wear* **2008**, 265 (3–4), 422–428.
- (7) Radice, S.; Mischler, S. *Wear* **2006**, 261 (9), 1031041.
- (8) Hu, X. L.; Gong, H. B.; Wang, Y. Z.; Chen, Q.; Zhang, J.; Zheng, S. H.; Yang, S. K.; Cao, B. Q. *J. Mater. Chem.* **2012**, 22, 15947–15952.
- (9) Zhou, G. H.; Zhu, Y. F.; Wang, X. M.; Xia, M. J.; Zhang, Y.; Ding, H. Y. *Wear* **2013**, 301 (1–2), 753–757.
- (10) Cao, S. W.; Zhu, Y. J.; Ma, M. Y.; Li, L.; Zhang, L. J. *J. Phys. Chem. C* **2008**, 112 (6), 1851–1856.
- (11) Peng, D. F.; Beyesen, S.; Li, Q.; Jian, J. K.; Sun, Y. F.; Jiwuer, J. L. *Particuology* **2009**, 7 (1), 35–38.
- (12) Visalakshi, G.; Venkateswaran, G.; Kulshreshtha, S. K.; Murphy, P. H. *Mater. Res. Bull.* **1993**, 28 (8), 829–836.
- (13) Pu, Z. F.; Cao, M. H.; Yang, J.; Huang, Y. L.; Hu, C. W. *Nanotechnology* **2006**, 17 (3), 799–804.
- (14) Woo, K. J.; Lee, H. J.; Ahnm, J. P.; Park, Y. S. *Adv. Mater.* **2003**, 15 (20), 1761–1764.
- (15) Lu, J.; Jiao, X. L.; Chen, D. R.; Li, W. J. *J. Phys. Chem. C* **2009**, 113 (10), 4012–4017.
- (16) Bondi, S. N.; Lackey, W. J.; Johnson, R. W.; Wang, X.; Wang, Z. L. *Carbon* **2006**, 44 (8), 1393–1403.
- (17) Dave, H. A.; Blank, G.; Rijnders, J. H.; Gertjan, M. K.; Rogalla, H. *Appl. Surf. Sci.* **1998**, 127, 633–638.
- (18) Fukata, N.; Oshima, T.; Tsurui, T.; Ito, S.; Murakami, K. *Sci. Technol. Adv. Mater.* **2005**, 6 (6), 628–632.
- (19) Munoz, E.; Maser, W. K.; Benito, A. M.; Martinez, M. T.; DelaFuente, G. F.; Maniette, Y.; Righi, A.; Anglaret, E.; J. Sauvajol, L. *Carbon* **2000**, 38 (10), 1445–1451.
- (20) Zeng, H. B.; Du, X. W.; Singh, S. C.; Kulinich, S. A.; Yang, S. K.; He, J. P.; Cai, W. P. *Adv. Funct. Mater.* **2012**, 22, 1333–1353.
- (21) Yang, G. W. *Prog. Mater. Sci.* **2007**, 52 (4), 648–698.
- (22) Fojtik, A.; Henglein, A. B.; Buns, J. *Phys. Chem. C* **1993**, 97 (2), 252–254.
- (23) Liu, J.; Liang, C. H.; Xu, G. P.; Tian, Z. F.; Shao, G. S.; Zhang, L. D. *Nano Energy* **2013**, 2, 328–336.
- (24) Qin, W. J.; Sun, J.; Yang, J.; Du, X. W. *Mater. Chem. Phys.* **2011**, 130 (1–2), 425–430.
- (25) Hu, X. L.; Gong, H. B.; Xu, H. Y.; Wei, H. M.; Cao, B. Q.; Liu, G. Q.; Zeng, H. B.; Cai, W. P. *J. Am. Ceram. Soc.* **2011**, 94 (12), 4305–4309.
- (26) Zeng, H. B.; Cai, W. P.; Li, Y.; Hu, J. L.; Liu, P. S. *J. Phys. Chem. B* **2005**, 109 (39), 18260–18266.
- (27) Zeng, H. B.; Yang, S. K.; Cai, W. P. *J. Phys. Chem. C* **2011**, 115 (12), 5038–5043.
- (28) Yang, S. K.; Cai, W. P.; Zeng, H. B.; Xu, X. X. *J. Mater. Chem.* **2009**, 19, 7119–7123.
- (29) Yang, S. H.; Kiraly, B.; Wang, W. Y.; Shang, S. L.; Cao, B. Q.; Zeng, H. B.; Zhao, Y. H.; Li, W. Z.; Liu, Z. K.; Cai, W. P.; Huang, T. J. *Adv. Mater.* **2012**, 24, 5598–5603.
- (30) Yang, S. H.; Cai, W. P.; Zhang, H. W.; Xu, X. X.; Zeng, H. B. *J. Phys. Chem. C* **2009**, 113, 19091–19095.
- (31) Jayawardena, K. D. G. I.; Opoku, C.; Fryar, J.; Silva, S. R. P.; Henley, S. *Appl. Surf. Sci.* **2011**, 257 (12), 5274–5277.
- (32) Wang, H. Q.; Kawaguchi, K. J.; Pyatenko, A.; Li, X. Y.; Warkocka, Z. S.; Katou, Y.; Koshizaki, N. *Chem.—Eur. J.* **2012**, 18, 163–169.
- (33) Wang, H. Q.; Koshizaki, N.; Li, L.; Jia, L. C.; Kawaguchi, K. J.; Li, X. Y.; Alexander, P.; Zaneta, S. W.; Bando, Y.; Dmitri, G. *Adv. Mater.* **2011**, 23 (16), 1865–1870.
- (34) Ni, Y. H.; Ge, X. W.; Zhang, Z. C.; Ye, Q. *Chem. Mater.* **2002**, 14 (1), 1048–1052.
- (35) Zhang, D. H.; Liu, Z. Q.; Han, C.; Li, B.; Stewart, M. P.; Tour, J. M.; Zhou, C. W. *Nano Lett.* **2004**, 4, 2151–2155.
- (36) Fujii, T.; Groot, F. M. F.; Sawatzky, G. A.; Voogt, F. C.; Hibma, T.; Okada, K. *Phys. Rev. B* **1999**, 59, 3195–3202.
- (37) Liu, Q. X.; Yang, G. W.; Zhang, J. X. *Chem. Phys. Lett.* **2003**, 373 (1–2), 57–61.
- (38) Zhang, H. M.; Liang, C. H.; Tian, Z. F. *J. Phys. Chem. C* **2010**, 114 (29), 12524–12548.
- (39) Jozwiak, W. K.; Kaczmarek, E.; Maniecki, T. P.; Ignaczak, W.; Maniukiewicz, W. *Appl. Catal. A. Gen.* **2007**, 326 (1), 17–27.
- (40) Watanabe, Y. I.; Kuniyoshi, N.; Ishikawa, K.; Furuya, J.; Kato, M. *J. Phys.: Condens. Matter.* **2002**, 14 (49), 13643–13651.
- (41) Wimmers, O. J.; Arnoldy, P. J.; Moulijn, A. *J. Phys. Chem.* **1986**, 90 (7), 1331–1337.
- (42) Hernández Battez, A.; Viesca, J. L.; González, R.; Blanco, D.; Sedegbega, E. A.; Osorio, A. *Wear* **2010**, 268 (1–2), 325–328.
- (43) Shi, G.; Zhang, M. Q.; Rong, M. Z.; Bernd, W.; Klaus, F. *Wear* **2004**, 256 (11–12), 1072–1081.
- (44) Hernandez Battez, A.; Fernandez Rico, J. E.; Navas Arias, A.; Viesca, J. L.; Rodriguez, A.; Rodriguez, R. C.; Diaz Fernandez, J. M. *Wear* **2006**, 261 (3–4), 256–263.
- (45) Zhang, B. S.; Xu, B. S.; Xu, Y.; Gao, F.; Shi, P. J.; Wu, Y. X. *Tribol. Int.* **2011**, 44 (7–8), 878–886.
- (46) Kang, X. H.; Wang, B.; Zhu, L.; Zhu, H. *Wear* **2008**, 265 (1–2), 150–154.
- (47) Song, X. Y.; Zheng, S. H.; Zhang, J.; Li, W.; Chen, Q.; Cao, B. Q. *Mater. Res. Bull.* **2012**, 47 (12), 4305–4310.