

Quantized friction across ionic liquid thin films†

Cite this: *Phys. Chem. Chem. Phys.*, 2013, **15**, 15317
Alexander M. Smith,^a Kevin R. J. Lovelock,^b Nitya Nand Gosvami,^c Tom Welton^b
and Susan Perkin^{*a}

Received 3rd July 2013,
Accepted 6th August 2013

DOI: 10.1039/c3cp52779d

www.rsc.org/pccp

Ionic liquids – salts in the liquid state under ambient conditions – are of great interest as precision lubricants. Ionic liquids form layered structures at surfaces, yet it is not clear how this nano-structure relates to their lubrication properties. We measured the friction force between atomically smooth solid surfaces across ionic liquid films of controlled thickness in terms of the number of ion layers. Multiple friction-load regimes emerge, each corresponding to a different number of ion layers in the film. In contrast to molecular liquids, the friction coefficients differ for each layer due to their varying composition.

The importance of friction, and its reduction by lubrication of the sliding surfaces, has been recognized since ancient times and yet some of the most fundamental questions in this area remain unresolved.^{1,2} The classical laws of friction and lubrication do not apply at the nano-scale, where molecular interactions and arrangements determine the surface forces.^{2–4} The current renaissance in the study of friction mechanisms is motivated by the need to understand and control shear motion and lubrication in microscopic devices and between nano-scale objects.^{5–9} Of particular significance in microscopic systems, or when the external load is relatively small, is the adhesion contribution to the measured friction force.^{9–12} The presence of a thin liquid film between two solid surfaces can dramatically modify the friction and adhesion forces between them, and a particularly interesting case arises when the liquid molecules are arranged in layers between the surfaces leading to multiple adhesion values corresponding to different numbers of ion layers in the film.^{13–15}

Ionic liquids¹⁶ are a particularly attractive class of fluids for precision tribological applications due to their low melting point and wide liquid range, extremely low volatility, high

thermal stability, lubrication and wear-protection characteristics, and electrical and thermal conductivity.^{17–19} The interfacial structure of ionic liquids at charged surfaces consists of alternating positive and negative ion layers extending several layers into the bulk.^{20,21} This alternating cation/anion structure is even more striking when the liquid is confined to a film between two surfaces: measurements of the surface interaction force as a function of surface separation reveal large oscillations, substantially stronger than van der Waals or mean-field electrostatic interactions, corresponding to sequential squeeze-out of layers from between the surfaces. In contrast to molecular liquids, ionic liquids squeeze out two layers at a time – one cation layer and one anion layer – in order to maintain electroneutrality in the film.^{15,21–25} Despite the burgeoning interest in ionic liquids and their intricate nano-structure, very little is currently known about their mechanism of lubrication.^{26,27} To address this, we performed measurements of friction across films of ionic liquid with key features distinct from earlier studies^{15,26–28} and allowing new quantitative analysis: friction is measured across ionic liquid films of precisely controlled thickness (number of ion layers), at both positive and negative applied loads, and the adhesion and real contact area are measured concurrently. Our high resolution measurements reveal the following notable features for the first time in ionic liquid: (i) discrete friction regimes for each integer number of ion layers in the film, leading to discrete multi-valued ('quantized') friction as a function of load; (ii) stick-slip friction observed below a critical velocity (different for each number of ion layers); and (iii) different friction coefficients measured for each film thickness. This latter point implies that the mechanism of friction across ionic liquids is qualitatively different to non-polar liquids, where the friction coefficient is independent of film thickness.¹⁴ Put together, these findings point at new ways to control friction in microscopic systems and to develop ionic liquid lubricants for specialized applications.

The ionic liquid, 1-butyl-1-methylpyrrolidinium bis[(trifluoromethane)sulfonyl]imide or [C₄C₁Pyrr][NTf₂] (Fig. 1(a)), was confined between two mica sheets in a Surface Force Balance (SFB) (Fig. 1(b), and further details provided in ESI†). The mica sheets are each ~1 cm² area and ~2 μm thickness, atomically smooth and

^a Department of Chemistry, Physical and Theoretical Chemistry Laboratory, University of Oxford, South Parks Road, Oxford OX1 3QZ, UK.
E-mail: susan.perkin@chem.ox.ac.uk

^b Department of Chemistry, Imperial College London, London SW7 2AZ, UK

^c Department of Mechanical Engineering and Applied Mechanics, University of Pennsylvania, 220 S. 33rd St., Philadelphia, PA 19104, USA

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c3cp52779d



Fig. 1 Normal and shear forces between mica sheets across films of [C₄C₁Pyr][NTf₂]. (a) The cation and anion structures for the ionic liquid [C₄C₁Pyr][NTf₂]. (b) Schematic showing the system of ionic liquid arranged in layers between negative mica surfaces. (c) F_N vs. D showing the oscillatory force law due to sequential squeeze-out of pairs of ion layers, labeled with the number of ion layers at each distance and the corresponding values of F_{adh}/R ; data reproduced with permission.²⁴ (d) $F_{S,k}$ vs. F_N measured for different numbers of ion layers, i , in the film. The lines are linear fits to the data according to eqn (1). Inset diagrams indicate the liquid structure for each regime, and the circled data points are those corresponding to the traces in (e). (e) Friction loops showing F_L vs. lateral displacement, x_s , showing three examples measured at similar F_N and v_s but different numbers of ion layers, demonstrating the discrete or quantized nature of the friction ($i = 9, 7$, and 5).

step-free over the whole area, mounted in crossed-cylinder orientation with radius of curvature $R \sim 0.01$ m. When immersed in ionic liquids, mica surfaces have a high negative charge due to dissociation of K⁺ ions from the lattice.^{15,29} The thickness of the liquid film between the sheets was measured with sub-molecular resolution using optical interference according to a method described in detail earlier.³⁰ We measured the normal interaction force, F_N , as a function of mica-mica surface separation, D , and the lateral force, F_L , between them (as a function of D and F_N) which arises when one surface is translated by a lateral displacement x_s at velocity v_s . The measured friction responses contained certain features which are defined here as follows (and indicated in Fig. 2): If the film supports a finite stress then during initial translation F_L will increase up to a *yield* force, $F_{S,y}$. If shearing then proceeds by smooth sliding, the lateral force required to maintain sliding is the *kinetic friction* force, $F_{S,k}$. If shearing proceeds by stick-slip, F_L ranges between a minimum of $F_{S,k}$ at the stick-point and a maximum of the *static friction* force, $F_{S,st}$, at the slip-point. If $F_{S,y} > F_{S,k}$ (or $F_{S,y} > F_{S,st}$ in the case of stick-slip) then a *stiction spike* is observed, of height $F_{S,y} - F_{S,k}$.

The profile of F_N with D is an oscillatory function (Fig. 1(c)) due to the packing of the ions into layers and the sequential squeeze-out of pairs of ion layers, as expected for ionic liquids:

in the case of [C₄C₁Pyr][NTf₂] between negatively charged surfaces the stable film structures contain odd-integer numbers of ion layers,²⁴ which we label as $i = 3, 5, 7$, or 9 , as shown schematically in the insets to Fig. 1(d). As a result of the oscillatory force-law between the surfaces across the liquid it is possible to vary F_N – over a certain range including positive and negative values – while the number of ion layers, i , in the film remains constant. By performing this compression-decompression at constant i , and at the same time applying lateral displacement to the top surface and detecting the resulting F_L , we were able to measure $F_{S,k}$ as a function of F_N for each number of ion layers between the surfaces. The result reveals quantized friction regimes (Fig. 1(d)). Thus there exists a friction-load relationship for each value of i , and the friction is multi-valued for each single value of load. This latter point is illustrated in Fig. 1(e) where closely similar values of F_N give rise to different values of $F_{S,y}$ and $F_{S,k}$ depending on the number of ion layers.

Each of the quantized friction regimes follows an approximately linear $F_{S,k}$ vs. F_N relationship with non-zero $F_{S,k}$ measured at zero (and negative) values of F_N . The friction at zero applied load originates in the contribution from the adhesion force between the surfaces, F_{adh} , which is different for each i , as seen



Fig. 2 Friction loops, showing F_L as a function of x_s at a constant film thickness of $i = 7$ and for three different values of F_N . The velocity is constant at $v_s = 205 \text{ nm s}^{-1}$, and the back-and-forth displacement is in the directions indicated by (red) dashed arrows. Friction features $F_{S,y}$, $F_{S,k}$, and $F_{S,st}$ are defined for the $F_N = 160.4 \text{ μN}$ trace.

from the F_N minima in Fig. 1c. $F_{S,k}$ can be expressed as a linear combination of load-dependent and adhesion-dependent contributions:¹⁰

$$F_{S,k}^i = \mu^i F_N + \alpha^i \left(\frac{A^i \cdot F_{adh}^i}{\pi d^i R} \right) \quad (1)$$

where μ is the load-controlled friction coefficient, α is the adhesion-controlled friction coefficient, A^i is the flat contact

area, d^i is the distance between the shearing liquid layers, and superscripts i denote values for a film containing i layers. In our experiments we directly measured F_{adh}^i , $F_{S,k}^i$, R , and d^i whilst controlling F_N . A^i is related to F_N and $F_{S,k}^i$ by the JKR equation and was determined after *in situ* calibration of the elastic modulus.¹⁴ Thus it was possible to determine values of the coefficients μ^i and α^i for each discrete film thickness. We found that for the films consisting of 9, 7 and 5 ion layers μ^i increases gradually in the range 0.082–0.172 (± 0.008), whereas for $i = 3$ the gradient is much steeper giving $\mu^{i=3} = 0.812 \pm 0.090$. The adhesion contribution to the friction gives coefficients with closely similar values for all i ; $\alpha^i = 0.19 \pm 0.04$ (all values are provided in ESI†). The increase in μ^i as i decreases is likely due to the greater degree of interlocking of the layers for small i : the layers closer to the mica surfaces are more ordered, and have greater anion/cation excess concentration, and therefore have greater inter-layer attractions for the same value of contact area and a greater activation barrier for ‘unlocking’ the surfaces to allow shear.

To obtain further insight into the molecular reorganization taking place during shear we recorded high resolution traces of F_L as a function of lateral displacement of the top surface for a range of F_N and v_s . Fig. 2 and 3 show examples for $i = 7$; qualitatively similar results were observed for $i = 9, 7, 5$, and 3. Over the range of F_N and v_s studied the films exhibit a well defined yield point, with $F_{S,y}$ increasing in magnitude as F_N increases, and stiction spikes at the onset of sliding (Fig. 2). At low v_s , shearing of the film then progresses by way of a series of stick-slip cycles characterized by saw-tooth traces (Fig. 3(a)). At a critical v_s (Fig. 3(b)) the stick-slip disappears and above this velocity the film shears with a smooth sliding motion (Fig. 3(c)). It was notable that $F_{S,k}$ is independent of v_s , whilst $F_{S,y}$ is weakly dependent on v_s . Once the direction of applied lateral motion is reversed, similar behavior is observed in the opposite direction.

The clear yield points indicate solid-like behavior of the ionic liquid confined to films of $i \leq 9$ ion layers. Subsequent stick-slip behavior observed at lower v_s is most often attributed to a series of freeze-melt transitions.^{5,31} However in the case of



Fig. 3 Friction loops for varying v_s at constant F_N . (a) $v_s = 205 \text{ nm s}^{-1}$; (b) $v_s = 512 \text{ nm s}^{-1}$; (c) $v_s = 1024 \text{ nm s}^{-1}$; each measured at $F_N = 52.6 \text{ μN}$. $F_{S,y} > F_{S,k}$ leading to a stiction spike in all cases, but only at slower v_s are regular stick-slip cycles observed. At the critical velocity, here 512 nm s^{-1} , one stick-slip cycle is followed by smooth sliding. At higher v_s smooth sliding occurs after the initial spike.



ionic liquids a full-film melt is unlikely due to the high Coulombic energy barrier of ions swapping between layers, so the stick-slip is likely to involve either interlayer slips³² or intra-layer (two dimensional) melting. Furthermore, the particularly high values of $F_{S,k}^{i=3}$ compared to $F_{S,k}^{i>3}$ indicates that mid-film cation layers – the structural feature present in films of greater thickness but not in the $i = 3$ film – may shear-melt or slip at lower stress than the other layers and thus be responsible for the lower yield force of the thicker films.

The quantized friction presented here is a direct result of the multiple discrete values of the adhesion-controlled contribution to the total friction, and is expected to be general for liquids which form layered structures in thin films. Indeed, experiments in our laboratory with a range of other ionic liquids have revealed quantized regimes corresponding to the layer structure in every case. The result rationalizes previous reports of discontinuities in the friction across molecular liquids when the number of layers changes,^{10,14} here demonstrating multiple quantized friction regimes with varying friction coefficients and quantifying the effects of adhesion on the overall friction force. This result will be pivotal in explaining the boundary lubrication of rough surfaces where the liquid film is of varying thicknesses across the contact zone: the total friction is made up of differently weighted contributions from each film thickness despite uniform applied load. Thus the quantized friction regimes resolved here using atomically-smooth surfaces will provide a link between single-asperity and rough-contact friction.

Acknowledgements

This work was supported by The Leverhulme Trust (F/07 134/DK and F/07 134/DN), Taiho Kogyo Tribology Research Foundation, Weizmann UK, Infineum UK, The Office of Naval Research (N00014-10-1-0096) and the EPSRC (EP/J015202/1). Andrew Dolan is thanked for synthesis of the ionic liquid used in this work.

Notes and references

- 1 B. N. J. Persson, *Sliding Friction*, Springer-Verlag, 1998.
- 2 M. Urbakh, J. Klafter, D. Gourdon and J. Israelachvili, *Nature*, 2004, **430**, 525–528.
- 3 B. Bhushan, J. Israelachvili and U. Landman, *Nature*, 1995, **374**, 607–616.
- 4 Y. Mo, K. T. Turner and I. Szlufarska, *Nature*, 2009, **457**, 1116–1119.
- 5 P. A. Thompson and M. O. Robbins, *Science*, 1990, **250**, 792–794.
- 6 J. Y. Park, D. F. Ogletree, P. A. Thiel and M. Salmeron, *Science*, 2006, **313**, 186.
- 7 A. Socoliuc, E. Gnecco, S. Maier, O. Pfeiffer, A. Baratoff, R. Bennewitz and E. Meyer, *Science*, 2006, **313**, 207–210.
- 8 C. Lee, Q. Li, W. Kalb, X.-Z. Liu, H. Berger, R. W. Carpick and J. Hone, *Science*, 2010, **328**, 76–80.
- 9 R. Guerra, U. Tartaglino, A. Vanossi and E. Tosatti, *Nat. Mater.*, 2010, **9**, 634–637.
- 10 J. Israelachvili, Y. L. Chen and H. Yoshizawa, *J. Adhes. Sci. Technol.*, 1994, **8**, 1231–1249.
- 11 Z. Deng, A. Smolyanitsky, Q. Li, X.-Q. Feng and R. J. Cannara, *Nat. Mater.*, 2012, **11**, 1032–1037.
- 12 B. Luan and M. O. Robbins, *Nature*, 2005, **435**, 929–932.
- 13 R. Horn and J. Israelachvili, *J. Chem. Phys.*, 1981, **75**, 1400–1411.
- 14 E. Kumacheva and J. Klein, *J. Chem. Phys.*, 1998, **108**, 7010–7022.
- 15 S. Perkin, T. Albrecht and J. Klein, *Phys. Chem. Chem. Phys.*, 2010, **12**, 1243–1247.
- 16 J. P. Hallett and T. Welton, *Chem. Rev.*, 2011, **111**, 3508–3576.
- 17 M. Palacio and B. Bhushan, *Tribol. Lett.*, 2010, **40**, 247–268.
- 18 F. Zhou, Y. Liang and W. Liu, *Chem. Soc. Rev.*, 2009, **38**, 2590–2599.
- 19 A. Somers, P. Howlett, D. MacFarlane and M. Forsyth, *Lubricants*, 2013, **1**, 3–21.
- 20 M. Mezger, H. Schröder, H. Reichert, S. Schramm, J. S. Okasinski, S. Schöder, V. Honkumaki, M. Deutsch, B. M. Ocko, J. Ralston, M. Rohwerder, M. Stratmann and H. Dosch, *Science*, 2008, **322**, 424–428.
- 21 R. Atkin and G. G. Warr, *J. Phys. Chem. C*, 2007, **111**, 5162–5168.
- 22 K. Ueno, M. Kasuya, M. Watanabe, M. Mizukami and K. Kurihara, *Phys. Chem. Chem. Phys.*, 2010, **12**, 4066–4071.
- 23 I. Bou-Malham and L. Bureau, *Soft Matter*, 2010, **6**, 4062–4065.
- 24 A. M. Smith, K. R. J. Lovelock, N. N. Gosvami, P. Licence, A. Dolan, T. Welton and S. Perkin, *J. Phys. Chem. Lett.*, 2013, **4**, 378–382.
- 25 S. Perkin, L. Crowhurst, H. Niedermeyer, T. Welton, N. N. Gosvami and A. M. Smith, *Chem. Commun.*, 2011, **47**, 6572–6574.
- 26 J. Sweeney, F. Hausen, R. Hayes, G. B. Webber, F. Endres, M. W. Rutland, R. Bennewitz and R. Atkin, *Phys. Rev. Lett.*, 2012, **109**, 155502.
- 27 O. Werzer, E. D. Cranston, G. G. Warr, R. Atkin and M. W. Rutland, *Phys. Chem. Chem. Phys.*, 2012, **24**, 5147–5152.
- 28 A. Elbourne, J. Sweeney, G. B. Webber, E. J. Wanless, G. G. Warr, M. W. Rutland and R. Atkin, *Chem. Commun.*, 2013, **49**, 6797–6799.
- 29 A. M. Smith, K. R. J. Lovelock and S. Perkin, *Faraday Discuss.*, DOI: 10.1039/C3FD00075C.
- 30 S. Perkin, L. Chai, N. Kampf, U. Raviv, W. Briscoe, I. Dunlop, S. Titmuss, M. Seo, E. Kumacheva and J. Klein, *Langmuir*, 2006, **22**, 6142–6152.
- 31 M. Schoen, C. L. Rhykerd, D. J. Diestler and J. H. Cushman, *Science*, 1989, **245**, 1223–1225.
- 32 Y. Lei and Y. Leng, *Phys. Rev. Lett.*, 2011, **107**, 147801.

