Evaluation of Adhesion and Hysteresis Friction of Rubber–Pavement System

Mohammad Al-Assi * and Emad Kassem

Department of Civil and Environmental Engineering, University of Idaho, Moscow, ID 83844, USA; ekassem@uidaho.edu
* Correspondence: alas9935@vandals.uidaho.edu; Tel.: +1-208-310-4850

Received: 1 September 2017; Accepted: 30 September 2017; Published: 7 October 2017

Abstract: Tire-pavement friction is a key component in road safety. Adhesion and hysteresis are the two main mechanisms that affect the friction between rubber tires and pavements. This study experimentally examined the relationship between rubber–pavement adhesion and friction. The adhesive bond energy between rubber and pavement surfaces was calculated by measuring the surface energy components of rubber and aggregates. The friction was measured in the laboratory using a dynamic friction tester. The results revealed that there is a fair correlation between the adhesive bond energy and measured coefficient of friction. A rubber–pavement system with higher adhesion provided higher friction at low speed. In addition, the results demonstrated that there is a strong correlation between rubber–pavement friction and rubber properties. Softer rubber provided higher friction and vice versa. The results of this study provide an experimental verification of the relationship between adhesion and pavement surface friction. The adhesive bond energy and rubber rheological properties could be incorporated in computational models to study tire-pavement friction in different conditions (e.g., speed and temperature).

Keywords: adhesion friction; hysteresis friction; adhesive bond energy; hot mix asphalt

1. Introduction

Tire-pavement friction is one of the main factors that contribute to road safety. An adequate level of friction between the vehicle tire and pavement surface reduces the number of crashes, especially in wet pavement conditions [1]; therefore, an appropriate level of traction between rubber tires and pavements is essential for safe driving. Rubber is the principal component of vehicle tires. It is an elastomer that exhibits unique physical and chemical properties. It has low Young’s modulus and high yield strain, as compared with other materials.

It is documented that rubber generates three different forms of friction: adhesion, deformation, and wear. Previous studies have shown that the adhesion force is the most influential component in rubber friction in dry conditions and low speed [2,3]. It is postulated that the adhesion between rubber tires and pavements is highly influenced by the surface free energy, which depends on the chemical composition of rubbers. Yet, to the best of the authors’ knowledge, no studies have been carried out to investigate the effect of surface free energy of rubber on traction with pavements.

The theory of rubber friction recognizes three major friction forces: adhesion force, bulk deformations (hysteresis), and cohesion losses due to the wearing of rubber [4]. The adhesion friction force is formed due to rubber–pavement interaction at a microtexture level and highly dependent on the true area of contact, and thus the adhesion friction is dominant for smooth contact at low speeds and dry conditions [3]. The hysteresis component of friction is caused by the bulk deformation of rubber material as it comes into contact with pavement asperities. Such deformation causes energy losses as rubber slides over the pavement surface. These energy losses are caused by the hysteretic
losses in the rubber material due to loading and unloading [3]. The third major component of friction (cohesion losses) is caused by the wearing of rubber as it slides over the pavement surface. Pavement texture has an important role in the friction components—a recent study showed that higher pavement microtexture and macrotexture result in higher frictional properties [5]. The study also showed that pavement surfaces initially had lower frictional properties due to the thin film of asphalt coating the aggregates at the surface. The friction increased as the binder film was removed by traffic, exposing the microtexture. Then, the friction decreased due to polishing and abrasion of the aggregates at the surface.

2. Objectives

The main goal of this study was to investigate the effect of adhesion between rubber and pavement surface on friction at low speed. In addition, we evaluated the effect of rubber rheological properties on friction with pavements. These objectives were achieved by conducting the following tasks:

1. Measure the surface energy components of different rubber and aggregate materials.
2. Calculate the adhesive bond energy between rubber materials and aggregates.
3. Measure the rheological properties of test rubber materials at different temperatures and loading frequencies.
4. Measure the coefficient of friction between rubber and hot mix asphalt (HMA) substrates.
5. Investigate the correlation between the calculated adhesive bond energy and the measured coefficient of friction between rubber and pavement surfaces.
6. Investigate the correlation between the rubber dynamic modulus and the coefficient of friction.

We measured the surface free energy of test rubber materials using a Sessile drop device, while the surface energy components of aggregates were measured using a universal sorption device (USD). The surface free energy of the rubber materials and aggregates were used to calculate the adhesion between the rubber materials and aggregates used in preparing the HMA test substrates. The friction between rubber sliders prepared using test rubber materials and HMA substrates was measured using the dynamic friction tester (DFT) in dry conditions at low speed. In addition, we measured the dynamic modulus of test rubber materials at different temperatures and loading frequencies using a dynamic mechanical analyzer (DMA).

3. Theory of Surface Energy

There are two major components contributing to pavement friction: adhesion and hysteresis. The adhesion component of pavement friction is dominant at low speeds in dry conditions, while the hysteresis component is dominant at higher speeds in wet conditions [6]. The adhesion between any materials is a function of their surface free energy components, where the adhesion force is developed at the molecular level [7,8]. The atoms at surface have lower level of bond energy compared to the atoms in bulk. As molecules in the material bulk are surrounded by other molecules from all sides, an external work must be applied to create a new surface area [7]. This work is known as the total surface free energy of the material and is denoted with Greek letter gamma (γ), the surface free energy is measured in units of ergs/cm² or mJ/m². The two most recognized theories to explain the surface free energy are: the two-component theory, and the acid-base theory. According to the two-component theory, the total surface free energy is caused by dispersion forces (e.g., Lifshitz-van der Waals forces) and specific forces (e.g., H-bonding). The total surface free energy is the sum of these two forces as shown in the Equation (1) [9,10].

\[
\gamma = \gamma_{\text{Dispersive}} + \gamma_{\text{Specific}}
\]

According to the acid-base theory, the total surface free energy for any material has three components; nonpolar component also known as Lifshitz-van der Waals (\(\gamma_{LW}\)), and two polar components: Lewis acid (\(\gamma^+\)), and Lewis base (\(\gamma^-\)) [11,12]. The total surface energy can be calculated from these three components as given in Equation (2).
\[ \gamma_{\text{Total}} = \gamma_{\text{LW}} + 2\sqrt{\gamma_+ \gamma_-} \]  

(2)

Several theories explain the adhesion between two materials by studying the interfacial forces between the materials [13]. The adhesion between rubber tires and pavement surface is one mechanism leading to friction and is dominant at low speeds. The adhesion between rubber and aggregate can be measured if the surface free energies of both materials are known. The adhesion between two materials (A and B) is a function of the surface free energy components of these two materials, as given in Equation (3) [7].

\[ W_{AB} = 2\sqrt{\gamma_A^{\text{LW}} \gamma_B^{\text{LW}}} + 2\sqrt{\gamma_A^+ \gamma_B^-} + 2\sqrt{\gamma_A^- \gamma_B^+} \]  

(3)

The amount of work required to separate the two materials at the interface in vacuum is referred as the adhesive bond energy (\( W_{AB} \)). The adhesive bond energy between rubber sample (subscript A) and an aggregate (subscript B) can be calculated using Equation (3), while the cohesive bond energy of a single material (e.g., rubber) (\( W_{AA} \)) can be calculated according to Equation (4) [7].

\[ W_{AA} = \gamma_A^{\text{LW}} + 2\sqrt{\gamma_A^{\text{LW}} \gamma_B^{\text{LW}}} \]  

(4)

3.1. Methods for Measuring Surface Energy

3.1.1. Sessile Drop Method

The Sessile drop method is commonly used to measure the surface free energy between a probe liquid and solid surface by means of measuring the contact angle between a droplet of a selected probe liquid and material surface [14]. The Sessile drop test measures the contact angle between a drop of liquid of known surface energy and a solid material, to calculate the surface free energy (SFE) of solid material surfaces. The Young’s equation (Equation (5)), defines the equilibrium at the three-phase contact of solid-liquid and gas [15].

\[ \gamma_{SV} = \gamma_{SL} + \gamma_{LV} \cos (\theta) \]  

(5)

where \( \theta \) is the contact angle between a solid surface and a drop of probe liquid. The contact angle is measured from a static image using a charged-coupled device (CCD) camera as shown in Figure 1. The Young equation assumes that the surface is chemically homogeneous and topographically smooth.

![Figure 1. Contact angle between liquid and solid surfaces.](image)

In the Sessile drop technique, a droplet of probe liquid is placed on the test surface using a micro-syringe. The Sessile drop device (Figure 2a) uses a CCD camera static image to measure
the contact angle between the used probe liquid of known surface energy and the rubber surface. The contact angle is the average value of the measured right and left contact angles (Figure 2b).

Figure 2. (a) Sessile drop device; (b) Sketch of Sessile drop contact angle measurement.

To determine the surface energy components of a material, three different probe liquids are used in the test. However, in order to minimize experimental error in calculating the surface energy components, Little and Bhsain [7] recommended using five different probe liquids (water, glycerol, ethylene glycol, formamide, and diiodomethane) with known surface energy components. Table 1 summarizes the surface energy components of the five probe liquids used in this study.

Table 1. Surface free energy components for selected probe liquids (ergs/cm²).

| Probe Liquid     | $\gamma^{LV}$ | $\gamma^+$ | $\gamma^-$ | $\gamma^{Total}$ |
|------------------|---------------|------------|------------|------------------|
| Water            | 21.80         | 25.50      | 25.50      | 72.80            |
| Glycerol         | 34.00         | 3.92       | 57.40      | 64.00            |
| Formamide        | 39.00         | 2.28       | 39.60      | 58.00            |
| Methylene Iodide | 50.80         | 0.00       | 0.00       | 50.80            |
| Ethylene Glycol  | 29.00         | 1.92       | 47.00      | 47.99            |

3.1.2. Universal Sorption Device

The surface free energy components of aggregates are often measured using the universal sorption device (USD). In this method, the aggregate particles are placed in a sealed cell under vacuum and controlled temperature. The USD uses a magnetic suspension balance to measure the mass of the
aggregate particle and probe vapor adsorbed on the aggregate surface when aggregates are subjected
different pressures of probe vapor. The relationship between the amount of probe vapor adsorbed
on aggregate surface and the partial pressure is called the adsorption isotherm which is used to
determine the spreading pressure ($\pi_e$) \[7\]. The spreading pressure ($\pi_e$) is a function of the surface
energy components of aggregate particles (S) and probe vapor (V), as given in Equation (6).

$$\pi_e + 2 \gamma_v^{\text{Tot}} = 2 \sqrt{\gamma_s^{\text{LW}} \gamma_v^{\text{LW}}} + 2 \sqrt{\gamma_s^{\text{LW}} \gamma_v} + 2 \sqrt{\gamma_s \gamma_v} \quad (6)$$

Several researchers used the USD to measure the surface free energy components of aggregates [16–18].
In this study, the USD test was used to determine the surface free energy of the gabbro and limestone
aggregates. The surface energy components of rubber and aggregates are used to calculate the adhesive
bond energy or work of adhesion ($W_{AB}$), (Equation (3)) between rubber sliders and HMA surfaces.

4. Test Materials and Laboratory Experiments

In this study, six different rubber materials were used to investigate the frictional properties
between laboratory-prepared rubber sliders and HMA slabs prepared from limestone and gabbro
aggregates. The rubber materials used in this study included pure gum, Styrene Butadiene Rubber
(SBR), Nitrile, Ethylene Propylene Diene Monomer (EPDM), Neoprene, and Butyl. These materials
were selected in testing as they cover a wide range of rubber properties in terms of dynamic modulus,
elongation, and durability. Table 2 summarizes the rubber properties as provided by the manufacturer.

| Rubber Type | Tensile Strength (PSI) | Elongation (%) | Composition         | Durometer      |
|-------------|------------------------|----------------|---------------------|----------------|
| Pure Gum    | 3000                   | 600            | Organic Gum         | 40 A Nominal   |
| SBR         | 800                    | 250            | Styrene Butadiene   | 65 Shore A     |
| EPDM        | 800                    | 400            | Synthetic           | 60 A Nominal   |
| Nitrile     | 950                    | 250            | Synthetic           | 50 A Nominal   |
| Neoprene    | 1000                   | 220            | Synthetic           | 60 A Nominal   |
| Butyl       | 1000                   | 350            | Isobutylene Isoprene| 55 Shore A     |

Two sources of aggregates were evaluated in this study (gabbro and limestone). The gabbro
and limestone aggregates were used to prepare HMA slabs. The gabbro is an igneous rock
while the limestone is a sedimentary rock. Gabbro has rough surface compared to limestone [19].
These aggregates are used in road construction in the State of Qatar. Table 3 presents the aggregate
gradation used to prepare both gabbro and limestone substrates. It should be noted that the same
aggregate gradation and binder type (Pen 60–70) were used in preparing the asphalt mixture substrates.
The optimum binder content was found to be 4.3% for gabbro and 4.7% for limestone. The mixing and
compaction temperatures were 143 °C and 135 °C, respectively.

| Sieve Size | % Passing |
|------------|-----------|
| 1.5”       | 100.0     |
| 1”         | 98.6      |
| 3/4”       | 88.2      |
| 1/2”       | 76.9      |
| 3/8”       | 68.9      |
| N4         | 47.1      |
| N8         | 26.5      |
| N16        | 15.8      |
| N30        | 10.5      |
| N50        | 7.9       |
| N100       | 6.1       |
| N200       | 4.2       |
| Pan        | 0.0       |
The asphalt mixtures were prepared following the AASHTO T-312, and the test slabs were compacted using a small vibratory compactor as shown in Figure 3.

5. Measuring Frictional Properties

We used a dynamic friction tester (DFT) to measure the coefficient of friction between rubber sliders (Figure 4), prepared using the test rubber materials, and HMA substrate at low speed (20 km/h) in dry conditions. The DFT uses three rubber sliders attached to a rotating desk (Figure 5). The rotating desk is lowered on the surface once the specified speed is reached and the coefficient of friction is measured with speed until the rotating desk comes to a complete stop.

Figure 3. Preparing hot mix asphalt (HMA) slabs.

Figure 4. Prepared rubber sliders from test rubber materials.
For the prepared HMA substrates, the aggregates on the surface were coated with a thin asphalt binder. We used a three-wheel polisher to remove this thin film of binder to ensure direct contact between rubber sliders and aggregates. The surface energy components of asphalt binder are different than those of aggregate samples and thus it may affect the adhesive bond energy between rubber and surface aggregates. In the field, this thin layer of asphalt binder is removed under traffic in a short period of time. The three-wheel polisher consists of pneumatic rubber wheels that rotate over the test HMA substrate as shown in Figure 6. The DFT device was used to measure the coefficient of friction between the prepared rubber sliders (Figure 4) and HMA substrates.

6. Measuring Rubber Properties

The dynamic mechanical analyzer (DMA) device was used to measure the dynamic modulus of rubber samples tested in this study (Figure 7). The rubber test sample for the DMA testing was 50 mm in length, 16.8 mm in width, and 6.4 mm thick. The test was performed at different test temperatures (0 to 70 °C) and frequencies (0.1–70 Hz). The temperature was controlled during the test by placing
the DMA inside an environmental chamber. The dynamic modulus is calculated by dividing the maximum applied cyclic tensile stress by the resulting strain.

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7. Test Results and Discussion

7.1. Adhesion Friction

The average contact angle between rubber samples and the test probe liquids was recorded using the Sessile drop device. Each measurement represents the average contact angle measured from right and left. The results for the measured contact angles in degrees are presented in Table 4. Based on the contact angle measurements, the surface energy components were calculated for each rubber material using Equation (7), which is discussed in detail by Little and Bhasin [7].

$$W_{LS} = \gamma_L (1 + \cos \theta) = 2\sqrt{\gamma^L_W \gamma^L_s} + 2\sqrt{\gamma^S_W \gamma^V_s} + 2\sqrt{\gamma^S_s \gamma^V_s}$$  \hspace{1cm} (7)

The surface free energy components are summarized in Table 5. The surface free energy components of the aggregates were measured using a Universal Sorption Device (USD) from a previous study [20]. The surface free energy components for the gabbro and limestone aggregates are shown in Table 6. It was found that gabbro rock has the higher total surface free energy compared to limestone.

| Probe Liquid | Water | Glycerol | Ethylene Glycol | Formamide | Diiodomethane |
|--------------|-------|----------|-----------------|-----------|---------------|
| Rubber Type  | Contact Angle | Std. Dev | Contact Angle | Std. Dev | Contact Angle | Std. Dev | Contact Angle | Std. Dev | Contact Angle | Std. Dev |
| Pure gum     | 89.47 | 0.69     | 120.18 | 0.86 | 87.14 | 0.74 | 83.03 | 1.21 | 67.43 | 0.76 |
| SBR          | 100.98 | 0.86 | 101.28 | 1.47 | 87.16 | 1.05 | 84.19 | 0.66 | 71.74 | 0.55 |
| Nitrile      | 89.04 | 0.74 | 94.77 | 0.82 | 84.32 | 0.89 | 90.35 | 0.99 | 63.03 | 1.04 |
| EPDM         | 108.36 | 0.74 | 112.45 | 0.67 | 89.61 | 0.75 | 93.29 | 0.87 | 72.22 | 13.12 |
| Neoprene     | 124.58 | 0.89 | 112.02 | 0.74 | 90.23 | 0.95 | 87.54 | 1.75 | 69.63 | 1.64 |
| Butyl        | 111.32 | 0.96 | 117.02 | 1.21 | 92.49 | 0.71 | 107.16 | 0.93 | 64.43 | 0.98 |
Table 5. Measured surface energy of rubber materials using the Sessile drop device.

| Rubber   | Surface Energy Components (ergs/cm²) | Standard Deviation (ergs/cm²) |
|----------|--------------------------------------|-----------------------------|
|          | $\gamma^{ LW}$ | $\gamma^+$ | $\gamma^-$ | $\gamma^{Total}$ | $\gamma^{ LW}$ | $\gamma^+$ | $\gamma^-$ |
| Pure gum | 21.21       | 0.00       | 2.26       | 21.21          | 0.44       | 0.00       | 0.42       |
| SBR      | 22.11       | 0.00       | 2.24       | 22.11          | 0.46       | 0.00       | 0.40       |
| Nitrile  | 17.08       | 0.00       | 10.33      | 17.08          | 0.54       | 0.00       | 0.67       |
| EPDM     | 16.77       | 0.00       | 0.35       | 16.77          | 1.53       | 0.00       | 0.17       |
| Neoprene | 15.30       | 0.00       | 0.00       | 15.30          | 0.69       | 0.00       | 0.00       |
| Butyl    | 18.79       | 0.00       | 0.00       | 18.79          | 0.51       | 0.00       | 0.00       |

Table 6. Measured surface energy of aggregates using universal sorption device (USD) device, (adapted from [20]).

| Material | Surface Energy Components (ergs/cm²) |            |
|----------|--------------------------------------|------------|
|          | $\gamma^{ LW}$ | $\gamma^+$ | $\gamma^-$ | $\gamma^{Total}$ |
| Limestone| 69.35       | 0.28       | 1075.40    | 104.18          |
| Gabbro   | 57.37       | 3.34       | 6277.96    | 346.85          |

Figure 8. Adhesive bond energy between rubbers and aggregates materials.

The adhesive bond energy between different rubber and aggregate samples was calculated from their surface free energy components according to Equation (3). Figure 8 shows the calculated adhesive bond energies between the rubber and aggregates. The SBR and Pure Gum rubber materials were found to provide the highest adhesion with test substrates made with gabbro and limestone. The adhesive bond energy between limestone and SBR rubber was 79.91 ergs/cm². The Neoprene was found to provide the lowest adhesion with HMA substrates compared to other rubber materials. Also, the limestone was found to provide higher adhesion with rubber materials compared to gabbro in most cases.
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![Figure 9a](image1.png)

![Figure 9b](image2.png)

**Figure 9.** Measured coefficient of friction vs. the adhesive bond energy for (a) limestone and gabbro slabs separately (b) both gabbro and limestone slabs.

The coefficient of friction between rubber sliders and HMA substrates was measured using the DFT at low speed (20 km/h) in dry conditions. The adhesion component of the friction is dominant at low speed. Figure 9a shows the relationship between the adhesive bond energy and the coefficient of friction for gabbro and limestone separately, while Figure 9b shows the relationship between the coefficient of friction versus the adhesive bond energy between test rubber materials and all the test HMA substrates (both gabbro and limestone). It was found that there is a fair correlation between the adhesive bond energy and measured coefficient of friction. This relationship demonstrates that higher adhesion between rubber and pavement surface increases the coefficient of friction at low speed as one expects; however, this study provided an experimental verification to such relationship. In the meantime, we believe that this relationship is affected by two experimental limitations. First, the asphalt binder film was not fully removed from the aggregates on the surface. We attempted to minimize this effect by removing most of the film thickness by polishing the slab using the three-wheel polisher for 5000 cycles until the aggregate particles surface is exposed. The application of more cycles was found to polish the aggregates which should be avoided since the surface energy components of aggregate will change with change in the texture of the surface. Second, the surface energy components of rubber were calculated based on the contact angle between smooth rubber samples and probe liquids.
Meanwhile, we observed that the surface of rubber sliders, used in DFT, was not completely smooth due to cutting the rubber sheet to prepare the sliders. These two limitations may contribute to this fair correlation between adhesive bond energy and measured coefficient of friction at low speed.

7.2. Hysteresis Friction

The DMA results demonstrated that the dynamic modulus (E*) of rubber material increased with the loading frequency and decreased with temperature as expected. Figure 10 shows the E* master curves of the test rubber materials. The Pure Gum rubber provided the lowest E* value compared to other rubber materials, while the Nitrile rubber had the highest E*. From Figure 10, it can be clearly seen that the Pure Gum rubber is the softest followed by the SBR rubber while the Nitrile rubber is the stiffest. The rubber's dynamic modulus at room temperature (20 °C) were correlated with the measured coefficient of friction.

![Figure 10. E* Master curves of the test rubber materials.](image)

Figure 10 shows the relationship between the dynamic modulus values of rubber samples and the coefficient of friction between rubber sliders and HMA substrates. There is a strong correlation between the complex modulus at room temperature (20 °C) and the coefficient of friction at 30 km/h. The researchers observed that there was no significant change in the coefficient of friction measured using DFT with speed after 30 km/h. Figure 11 demonstrates that softer rubbers provide higher coefficient of friction while stiffer rubbers provided lower coefficient of friction regardless the aggregate type.
8. Summary and Conclusions

This study experimentally examined the relationship between the rubber–pavement adhesion and friction. We calculated the surface energy components for rubber materials by measuring the contact angle between rubber materials and different probe liquids. The contact angle was measured using a Sessile drop device. Similarly, the surface energy components of aggregate samples were measured using a USD device. The adhesive bond energy between rubber and aggregates was calculated based on the surface energy components of both materials. In addition, we measured the coefficient of friction between different rubber materials and HMA substrates prepared using different rock types. The results showed that there is a fair correlation between adhesive bond energy and friction between rubber and pavement surfaces. In addition, the results demonstrated that there is a strong relationship between friction and rubber elastic modulus. Softer rubber provided higher friction. The results of this study provide an experimental verification of the relationship between adhesion and friction of rubber and pavement surface. The adhesive bond energy and rubber rheological properties could be incorporated in computational models to study tire-pavement friction in different conditions (e.g., speed and temperature). We recommend testing more aggregate types and rubber materials to evaluate the adhesion friction.

Acknowledgments: This publication was made possible by an NPRP award (NPRP No. 7-482-2-184: Thermo-Mechanical Tire-Pavement Interaction: Computational Modelling and Field Measurements) from the Qatar National Research Fund (QNRF – a member of The Qatar Foundation). The statements made herein are solely the responsibility of the authors.

Author Contributions: Mohammad Al-Assi and Emad Kassem conceived, designed and performed the experiments, analyzed the data, and wrote the paper.

Conflicts of Interest: The authors declare no conflict of interest.

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