Particle-laden Tubeless Siphon

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A tubeless siphon was created by sucking a 1% aqueous Polyox§ solution laden with particles from a beaker into a cylinder by a moving piston. The piston speed and particle concentration were varied. At very high rates of withdrawal, all the liquid could be removed before the siphon broke. In this case, the beaker was completely cleaned without a trace of liquid. The addition of small concentrations of small, nearly neutrally buoyant particles greatly enhanced the pulling power of the liquid, reducing the threshold speed of withdrawal at which the beaker was completely cleaned. At speeds of withdrawal smaller than the threshold not all of the fluid-particle mixture is pulled out of the beaker. The amount pulled out first increases, then decreases as the particle concentration is increased.

1. Introduction

The tubeless siphon is a well-known special effect found in polymeric liquids and not in Newtonian liquids. This effect is well documented and is described in most of the standard works on rheological fluid mechanics, i.e., Bird, Armstrong & Hassager (1977), Joseph (1990), and Macosko (1994). The effect may be described as follows: a fluid is sucked through a nozzle with the nozzle elevated above the surface of the liquid, as shown in figure 1. Instead of the liquid breaking as in a Newtonian liquid, an unsupported fluid column is drawn from a pool below into the nozzle above without breaking, as shown in figure 2.

Boonacker (1999) studied the screening properties of particle-laden viscoelastic fluids, using the apparatus, fluid and particles used in this study. She reported that when the suction speed was sufficiently high, only the fluid would enter the tube; the particles were filtered out as in the Uepler effect. This filtering is called a screening effect. The tubeless siphon with particles was observed in her experiments.

Here we do a systematic study of the tubeless siphon with particles, pointing to the previously undocumented “cleaning property” which may have applications for decontamination. At a threshold withdrawal speed of the piston, the tubeless siphon laden with particles leads to complete removal of all fluid and particles from the pool (figure 2a). If there are no particles in the fluid, or the concentration of particles is lower than a critical value, the fluid column breaks during siphoning leaving a measurable volume of liquid behind (figure 2b).

† Movies and photos of the experiments reported in this paper can be viewed at http://www.aem.umn.edu/research/particles/siphon/.
§ Polyox is a registered trademark of Union Carbide.
2. Experimental conditions

The apparatus used in the experiment is shown schematically in figure 1; it consists of three primary parts: a piston pump, a tube and a beaker. Pumping is done in a 8.89 cm long cylinder with 10.08 cm inner diameter and a piston driven by an Instron tensile machine. A tube with 1.27 cm outer diameter and 0.470 cm inner diameter is inserted into the cylinder from the bottom. The beaker serves as the reservoir for the viscoelastic fluid with particles.

The viscoelastic fluid used in the experiments was 1% by weight aqueous WSR 301 Polyox solution. Polyethene oxide has extremely long molecular chains with a molecular weight of 4,000,000 g/mol, giving non-Newtonian and viscoelastic characteristics to the de-ionized water in which it has been dissolved. The process of dissolving Polyox powder into water was slow and it took about one week of continuous mixing to obtain one gallon of homogeneous solution.

Two kinds of particles were used in the experiments. Spherical, colorless, polymeric beads were used in most runs. The density of the beads was 1.0 g/cc and the size was in the range of 850–1000 μm. Spherical brown resin particles with sizes in the range of
600–700 μm were also used; the brown particles are visible in photographs. The resin particles were nearly neutrally buoyant in 1% aqueous Polyox solution.

The following procedures were followed in the experiments. A well-mixed testing fluid with suspended particles was prepared. At the beginning of each run, the piston was at its lowest position in the cylinder. The beaker was lifted so that the tube would dip into the pool of liquid. The Instron tensile machine was started raising the piston at a constant speed. The pressure difference between the atmosphere and the vacuum in the cylinder drove the fluid into the cylinder through the tube. Once the siphon started, the beaker was lowered so that the nozzle of the tube was higher than the fluid surface and the tubeless siphon formed. As the piston rose at the constant speed, the tubeless siphon could either suck all the liquid out of the beaker, or the siphon would break, leaving behind a measurable volume of liquid. The volume of the fluid which entered the cylinder was also recorded.

The distance from the nozzle to the surface of the pool at the moment when the tubeless siphon started was kept at 15 cm (with error less than 1 cm). The temperatures under which the experiments were conducted were in the range of 21.1 °C to 22.78 °C. Several key rheological parameters of 1% aqueous Polyox solution can be found in Liu (1995). However, extensional viscosities are not available.

3. Analysis

To investigate the properties of the particle-laden tubeless siphon, it is necessary to understand the mechanism of the breaking of the tubeless siphon. The volume flow rate in the cylinder and the tube is given by,

\[ UA = V_t A_t \]  \hspace{1cm} (3.1)

where \( U \) is the speed of the piston and \( A \) is the inner cross sectional area of the cylinder; \( V_t \) is the velocity of the fluid in the tube and \( A_t \) is the inner cross sectional area of the tube (see figure 1). In each run, the piston speed \( U \) was kept constant giving rise to a constant \( V_t \); conservation of mass implies that

\[ V_t A_1 = V_0 A_0 \]  \hspace{1cm} (3.2)

where \( V_t \) and \( A_1 \) are the velocity and the cross sectional area of the fluid column at the nozzle and \( V_0 \) and \( A_0 \) are the velocity and the cross sectional area of the fluid column at the surface of the pool (figure 1).

Before the siphon breaks, \( V_t \) and \( A_1 \) are balanced by \( V_t \) and \( A_t \). Therefore, during the process of siphoning, the volume flux of the fluid drawn out of the pool (\( V_0 A_0 \)) is a constant. If the piston speed is high, it leads to a high extensional rate and high extensional force which can maintain a constant volume flux in the siphon until all the fluid is sucked out of the beaker. However, if the piston speed is low, the volume flux will

\[ \begin{array}{|c|c|c|c|c|}
\hline
T(°C) & \rho(g/cc) & \eta_p(Pa.s) & \psi_{10}(g/cm) & \lambda_0(s) \\
\hline
24 & 1 & 7.65 & 360 & 0.390 \\
\hline
\end{array} \]

Table 1. Key rheological parameters of 1% aqueous Polyox solution at 24 °C. \( T \) is the temperature; \( \rho \) is the density; \( \eta_p \) is the shear viscosity at zero shear rate; \( \psi_{10} \) is the first normal stress coefficient at zero shear rate; \( \lambda_0 \) is the longest relaxation time.
decrease at some point in the siphoning process, thereby

\[ V_0A_0 = V_1A_1 < V_tA_t \]  \hspace{1cm} (3.3)

At that moment, \( V_1 \) is still equal to \( V_t \), thereby \( A_1 < A_t \). This inequality implies that air will be sucked into the tube and the pressure difference between the atmosphere and the inside of the cylinder will vanish rapidly. Consequently, the tubeless siphon stops.

From the analysis and experiments, it is found that the cleaning properties depend on a number of conditions: the piston speed, the concentration of particles in the fluid, the properties of the fluid, the properties of the particles, the size and the shape of tube, and the distance of the nozzle from the pool. Moreover, temperature plays an important role because the properties of the Polyox solution are very sensitive to the temperature. This study will focus on the piston speed and the concentration of the particles in the fluid. The study of other conditions will be conducted in the future.

4. Experimental results

First we give results for the siphon of pure fluid and then for the siphon of the fluid-particle mixture.

4.1. Pure fluid siphon

The effect of the piston speed was studied in a series of runs using Polyox solutions without particles. In all these runs, the original volume of the Polyox solution in the beaker was 150 ml. The piston speed ranged from 0.0704 cm/s to 0.205 cm/s. When the siphon stopped, the volume of the Polyox solution left in the beaker was recorded (the error was \( \pm 5 \) ml). In table 2, the piston speed \( U \), the fluid temperature \( T \), and the suction fraction \( F \) (volume fraction sucked out of the original volume) are listed.

Table 2 indicates that the higher the piston speed, the more the fluid was sucked out. A higher piston speed leads to a higher extensional rate and larger extensional force which can pull more fluid out of the beaker. The threshold piston speed \( U_{Th} \) for pure fluid siphon under our experimental conditions is between 0.185 and 0.205 cm/s. The fluid is always completely cleaned when the piston speed is higher than the threshold.

When \( U \) was 0.0704 or 0.0895 cm/s, the volume of fluid sucked out was so small that it could not be measured reliably, hence, \( F \) was taken as zero in table 2 for \( U = 0.0704 \) and 0.0895 cm/s. If these two cases are excluded, the data can be described by a power law. \( F \) vs. \( U \) is plotted for \( U = 0.109, 0.128, 0.147, 0.166, 0.185 \) and 0.205 cm/s in figure 3; a power law fits the data extraordinarily well.

\[ F = 15600U^{5.94} \quad \text{for} \quad U < U_{Th} \]

\[ F = 1 \quad \text{for} \quad U \geq U_{Th} \]  \hspace{1cm} (4.1)

where 0.185 < \( U_{Th} < 0.205 \) cm/s.

Equation (4.1) shows that the piston speed has a large effect on the suction fraction
Figure 3. Power law fitting for $F$ vs. $U$. The fraction of fluid $F$ sucked out of the original volume increases with the piston speed $U$ raised to the power $5.94$. For $U \geq U_{Th}(0.185 < U_{Th} < 0.205 \text{ cm/s})$ all of the fluid is removed from the beaker; hence $F = 1$.

because the suction fraction is proportional to the piston speed raised to 5.94. The pre-factor and exponent in the power law correlation could be functions of other parameters, such as the polymer properties and the distance from the nozzle to the pool. The threshold piston speed for pure fluid siphon was lower when a Polyox solution with a molecular weight of $8,000,000$ g/mol instead of $4,000,000$ g/mol was used. We also believe that the threshold will be lower if a solution with higher Polyox concentration is used. Systematic tests of $U$ vs. $F$ for different liquids have not yet been done. It may be possible to use the beaker test as a kind of viscometer for extensional flows.

4.2. Particle-laden siphon

The concentration $C$ of particles is given as a percentage of the total volume:

$$C = \frac{\text{volume of particles}}{\text{volume of particles} + \text{volume of fluid}} \times 100$$  \hspace{1cm} (4.2)

Tests of the effects of concentration were done by varying the concentration in siphon experiments at a fixed speed. Particles were added to 150 ml of liquid to create increments of $C$. The mixture was stirred to be as homogenous as possible. To reduce the possibility of memory effects, the experiments were conducted at least 20 seconds after stirring. Since the longest relaxation time of the 1% aqueous Polyox solution is about 0.4 second (table 1), stirring should not have an appreciable effect on the test result. For each value of $C$ the volume left in the beaker after the siphon broke was recorded.

Experiments were carried out for $U = 0.0704, 0.0895, 0.109, 0.128, 0.147$ and $0.166 \text{ cm/s}$. The concentration of the particles ranged from 0.66% to 10%. In table 3, the piston speed $U$, the mixture temperature $T$, and the suction fraction $F$ pulled out of the original volume are listed.

The data in table 3 shows that for $U = 0.109, 0.128, 0.147$ and $0.166 \text{ cm/s}$, the presence of particles has dramatic affect on the suction fraction. At about 1% particle concentration, the mixture can be removed completely; without particles only a small fraction (less than 1/3) of the fluid can be sucked out. Studies of $F$ vs. $C$ for highly concentrated fluid-particle mixture were not done. Data in table 3 and figure 4 shows that $F$ first increases and then decreases with $C$ for $U = 0.0704$ and $0.0895 \text{ cm/s}$; complete cleaning was not observed at these speeds.

The threshold piston speed $U_{Th}$ at which a complete cleaning is obtained depends on the particle concentration. As shown in section 4.1, $0.185 < U_{Th} < 0.205 \text{ cm/s}$ for pure fluid siphon ($C = 0$). The presence of particles reduces the threshold value, i.e.,
The suction fraction \( F \) vs. the concentration of particles \( C \). Piston speed \( U = 0.0704 \), 0.0895, 0.109, 0.128, 0.147 and 0.166 cm/s.

\[
\begin{array}{cccc}
U(\text{cm/s}) & T(\text{°C}) & C(\%) & F \\
\hline
0.0704 & 21.83 & 0 & 0.000 \\
 & & 0.99 & 0.175 \\
 & & 1.96 & 0.183 \\
 & & 2.98 & 0.191 \\
 & & 4.00 & 0.200 \\
 & & 5.00 & 0.430 \\
 & & 6.02 & 0.248 \\
 & & 7.01 & 0.194 \\
 & & 10.02 & 0.190 \\
0.0895 & 22.61 & 0 & 0.000 \\
 & & 0.99 & 0.670 \\
 & & 1.96 & 0.869 \\
 & & 2.98 & 0.871 \\
 & & 4.00 & 0.712 \\
 & & 5.00 & 0.525 \\
 & & 7.01 & 0.225 \\
0.109 & 22.5 & 0 & 0.033 \\
 & & 0.99 & 0.967 \\
 & & 1.96 & 1.000 \\
0.128 & 22.44 & 0 & 0.067 \\
 & & 0.99 & 1.000 \\
 & & 1.96 & 1.000 \\
0.147 & 22.0 & 0 & 0.167 \\
 & & 0.99 & 1.00 \\
 & & 1.96 & 1.00 \\
 & & 10.02 & 1.00 \\
0.166 & 21.94 & 0 & 0.333 \\
 & & 0.66 & 1.00 \\
 & & 5.00 & 1.00 \\
\end{array}
\]

Table 3. The suction fraction \( F \) vs. the concentration of particles \( C \). Piston speed \( U = 0.0704 \), 0.0895, 0.109, 0.128, 0.147 and 0.166 cm/s.

Figure 4. The suction fraction \( F \) vs. the concentration of particles \( C \) < 12%. Piston speed \( U = 0.0704 \), 0.0895, and 0.147 cm/s. Data are from table 3. The plot denoted by \( U = 0.0704 \) is for the experiments at piston speed \( U = 0.0704 \) cm/s using a degraded Polyox solution stored in a refrigerator for six months.

\( U_T h(C > 0) < U_T h(C = 0) \). For example, table 3 shows that when \( C = 0.99 \), the threshold value reduces to 0.109 < \( U_T h < 0.128 \) cm/s. However, there exists a minimum threshold speed \( \bar{U}_T h \) so that when \( U < \bar{U}_T h \), complete cleaning cannot be achieved no matter what the particle concentration \( C \) is. Table 3 shows that the minimum threshold \( \bar{U}_T h \) is between 0.0895 and 0.109 cm/s in our experiments.

A critical concentration could be defined as the smallest concentration of particles at which the suction fraction \( F \) reach its peak value under a fixed \( U \). For \( U = 0.0704 \) and 0.0895 cm/s, the critical \( C \) is the \( C \) at which \( F \) peaks. For \( U = 0.109, 0.128, 0.147 \) and 0.166 cm/s, the critical \( C \) is the smallest \( C \) at which the complete removal is obtained.
For $U = 0.205$ cm/s, fluid without particles can be cleaned up. Hence, the critical $C$ is 0 for $U = 0.205$ cm/s. The critical $C$ for each value of $U$ are listed in table 4 and plotted in figure 5. The critical $C$ decreases as $U$ increases.

It is difficult to accurately determine the critical concentration mainly because the Polyox solution degrades very quickly after its creation and is also very sensitive to temperature. For these reasons, the precise values of the suction fraction are erratic and not easily reproduced. We did certain experiments to verify the existence rather than the precise value of a critical concentration of particles at piston speed below the threshold speed. To test the effects of degradation, we did experiments using two samples of the same Polyox solution; one was tested immediately after its creation and the other after six months of storage in a refrigerator. We believe that the refrigerated solution degraded by oxidation (see Bailey and Koleske (1976)). The $F$ vs. $C$ graph for the degraded solution is compared with the fresh solution at $U = 0.0704$ cm/s in figure 4. The critical concentration of particles for the degraded solution is higher than that for the fresh solution as would be expected; the maximum $F$ for the degraded solution is higher than that for the fresh solution probably because the testing temperature for the degraded solution is 0.9–1.2 °C lower than for the fresh solution. Another difference is that the $F$ vs. $C$ curve near the criticality for the fresh solution is more sharply peaked than for the degraded solution. It is probable that the $F$ vs. $C$ curves would be smooth in experiments in very finely divided intervals of $C$.

5. Mechanics of the particle-laden tubeless siphon

In this section, we propose some explanations of the mechanics of particle-laden tubeless siphon. When the tubeless siphon starts, the particles which are sucked up in the fluid column appear to be trapped in the free surface of the siphon, with few or no particles inside. Such a distribution of the particles is related to the velocity profile in the fluid column. According to Matthys (1988), the axial velocity at the centerline is much greater than the velocity of the free surface at the same axial location (see figure 6). The
velocity profile is similar to the velocity profile in the Poiseuille flow in a tube. However, there is no wall in the siphon flow, hence, no lubrication force to push the particles away from the free surface and strong return flow at the free surface driven by gravity can be observed. As a result, most of the particles are driven away from the centerline to the free surface.

Figure 7 shows a sequence of photos for one siphon experiment using brown resin particles. In figure 7(a) and 7(b), it can be seen that particles are brought up by the siphon but filtered out at the nozzle; almost none of them enter the tube. In experiments, we noted that particles screened from entering the tube fall in a return flow on the outside of the siphon. If the concentration of the particles is sufficiently high, the particles gather in the free surface of the fluid column, forming a pseudo-tube from the nozzle to the pool (see figure 2(a)). If the concentration of the particles is lower, the particles can not cover the free surface of the whole column. However, in the portion adjacent to the nozzle, due to the screening effects which will be discussed next, the surface of the column is almost always covered by particles (see figure 7). These particles apparently serve as an extension of the tube. The particles in the free surface could provide additional support for the tubeless siphon and improve the suction.

The screening effect reported in Boonacker (1999) plays an important role in the enhancement of the suction. The screening effect means that when the suction speed is sufficiently high, only the fluid enters the tube and the particles are filtered out. This effect is closely related to the characteristics of the entrance flow of viscoelastic fluids (Binding & Walters 1988). Figure 8 illustrates such characteristics schematically in comparison with the injection flow of the Newtonian fluids. Suction and injection are asymmetric affects and the asymmetry has opposite polarity in Newtonian and viscoelastic fluids. In Newtonian fluids, injection is a long range effect associated with the propagation of vorticity created at the nozzle exit as shown in figure 8(a). The flow into a nozzle does not generate vorticity and the streamline are like potential sink flow (figure 8(b)). One familiar consequence of the asymmetry of suction and injection in Newtonian fluids is that it is easy to blow out a candle even from great distances but it is nearly impossible to suck it out. The circulations which develop in viscoelastic fluid are opposite to those in Newtonian fluid: all the eddies are reversed. The secondary motion which develops in suction draws fluid remotely into the nozzle as shown in figure 8(c). The circulation which develops when a viscoelastic fluid is injected from a nozzle into a reservoir (figure 8(d))

Figure 6. The velocity profile in the fluid column in tubeless siphon according to Matthys (1988). In the experiments a gravity driven return flow at the free surface makes the difference in velocity between centerline and free surface even greater.
Newtonian injection
Effects are long range

Newtonian suction
Effects are short range

Viscoelastic suction
Effects are long range

Viscoelastic injection
Effects are short range

Figure 7. Photos for siphon experiments using brown resin particles. (a) and (b) show that nearly all the particles are filtered out and do not enter the nozzle. (c) and (d) show a cluster of particles around the nozzle and that the siphon continues even though the fluid column has a smaller radius than the tube. From Boonacker (1999).

Figure 8. Asymmetry of suction and injection in Newtonian and viscoelastic fluid (just the opposite).

is opposite to that shown for suction (figure 8(c)): it inhibits long range effects injection and tends to transfer fluid sideways in motions similar to source flow with streamlines reversed.

The higher the suction speed, the stronger the circulation. In the siphon flow near the nozzle, the particles in the free surface of the fluid column are trapped in the circulation and can not enter the nozzle. As a result, more and more particles accumulate near the nozzle and cover the upper portion of the fluid column. Due to this accumulation, a cluster of particles can often be seen around the nozzle (see figures 7c and 7d). When the volume of fluid left in the beaker decreases, the fluid column will neck down at the nozzle so that the cross-sectional area of the liquid column $A_1$ is smaller than the cross-sectional
area of the tube $A_r$; air will be sucked in and the liquid column ruptures, stopping the siphon. However, the presence of the cluster of particles around the nozzle can prevent air from entering the nozzle. The fluid column shown in figures 7(c) and 7(d) has a smaller radius than the tube, but the siphon does not stop. Thus, the presence of the particles prolongs the time of the siphon process and helps the suction.

When $U = 0.0704$ and $0.0895$ cm/s, increasing the concentration of particles beyond the critical value of $C$ caused a decrease of the suction fraction $F$. The reason for this is not understood.

6. Conclusion

The new results reported here are:
(a) The pulling power of the tubeless siphon of a 1% aqueous Polyox solution is greatly increased by the suction speed. When there are no particles in the fluid, the suction fraction is proportional to the suction speed raised to the power 5.94.
(b) At sufficiently high rates of withdrawal all the liquid may be removed by the siphon, leaving behind a dry and clean beaker. We call this a cleaning property of the tubeless siphon.
(c) The pulling power of a siphon is dramatically increased by the presence of small particles in concentrations less than 10%. At high suction speeds (0.109, 0.128, 0.147 and 0.166 cm/s) the presence of particles leads to complete removal of the particle-laden fluid from the beaker. At low suction speeds (0.0704 and 0.0895 cm/s) the suction fraction peaks in a certain range of particle concentrations and then decreases.
(d) The cleaning property of the particle-laden siphon is here related to the screening of particles at the tube entrance.

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