

LABORATORY WORK ON TOTAL SUCTION CALIBRATION CURVE FOR WHATMAN #42 FILTER PAPER

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ABSTRACT

Soil suction is one of main parameter in unsaturated soil mechanics as its measurement is as very important as pore water pressure measurement for effective stress concept in saturated soil mechanics. Basically, the measurement of soil suction can be classified into 2 categories; direct and indirect method. In the first method, negative pore water pressure is measured directly to obtain matric suction, whereas in indirect method, suction is obtained indirectly by using salt solution and censor. Calibration is then required to establish the relationship between concentration of salt solution and censor. Filter paper method is considered as the simplest method and relatively low cost compare to other method. The studies conducted by many investigators indicate that calibration curve from one type to another type of filter paper is quite different. This paper presents the basic concepts of laboratory work on establishing total suction calibration curve for Wahtmann #42 filter paper. The work has been conducted using NaCl and KCl solutions with different molality to generate different suction. The result indicates that the higher molality of solution, the higher suction is generated. The calibration curve from KCl exhibits a bit higher compare to the one from NaCl. The difference of both curves most probably is due to the difference of equilibration period and suction source.

Keywords: Calibration curve, filter paper, suction, unsaturated soil

1. GENERAL

Soil suction is one of main parameter in unsaturated soil mechanics. Its measurement is as very important as pore water pressure measurement for effective stress concept in saturated soil mechanics (Houston et al. 1994).

Basically, the measurement of soil suction can be classified into 2 categories: direct and indirect method. In the first method, negative pore water pressure is measured directly to obtain matric suction. Tensiometer and null type pressure plate are examples of this method. In indirect method, suction is obtained indirectly by using salt solution and censor. Calibration is required to establish the relationship between concentration of salt solution and censor. This calibration test usually performed in equilibrium at particular temperature. Psychrometer, electric/thermal conductivity censor and filter paper methods are three examples of indirect method.

Recently, filter paper method is considered as the simplest method and relatively low cost compare to other method. This method is able to measure high range of suction (0-30000 KPa). The studies conducted by many investigators indicate that calibration curve from one type to another type of filter paper is quite different. Although filter paper is an industrial material which is produced under high technology control, calibration should be made for every batch of filter paper (Marinho and Oliveira, 2005).

This paper presents theoretical concept and laboratory practice for filter paper calibration curve for total suction. The result of this study will briefly be compared and discussed to other available curves.

2. MATRIC, OSMOTIC, AND TOTAL SUCTION

There are some definitions of soil suction. Suction may be defined as unit attractive force of the soil for water. It also has been defined as force needed to take away a soil water molecule from the soil into vapour phase (Bulut, 1996).

Commonly, soil suction is defined as free energy state of soil water. The free energy of the soil water can be measured in terms of the partial vapour pressure of the soil water (Fredlund and Rahardjo, 1993).

There are 3 kinds of soil suction: total suction, matric suction and osmotic suction. Matric suction is suction coming from capillary force, soil texture, and adsorption forces of soil particles. Houston (1994) has defined that matric suction is the affinity of soil has for water in the absence of any salt content gradient in the water. Osmotic suction is suction coming from salt concentration differences between one point and another point in the soil mass. Total suction is suction coming from matric suction and osmotic suction. Total suction, matric suction and osmotic suction can be written in mathematics expression as:

$$\Psi = (u_a - u_w) + \pi \quad (1)$$

Where

ψ = total suction

$(u_a - u_w)$ = matric suction

π = osmotic suction

Thermodynamics theory may be applied as basic concept for total suction measurement. In thermodynamics term, total suction can be considered as free energy state of soil pore water, which may be measured in term of partial pressure vapour of soil pore water at equilibrium (Houston et al. 1994; Leong et al., 2002; Likos and Lu 2002, Bulut and Wray 2005, Marinho and Oliveira 2005). Total suction (in KPa) above free surface/flat of pure water can be determined by using Kevin's Equation as:

$$\psi = \frac{RT}{V} \ln \left(\frac{P}{P_0} \right) \quad (2)$$

Where

R = universal gas constant (8.31432 J mol⁻¹K⁻¹)

T = Absolute temperature (273.16 + t°C)

V = molecular volume of water, volume of 1000 moles of liquid water (0.018 m³)

P = partial pressure of pore water vapour (kPa)

P₀ = saturation pressure of water vapour over a flat surface of pure water at the same temperature (kPa)

If P/P₀ is called relative humidity, than the equation (2) can be written as:

$$\psi = \frac{RT}{V} \ln(RH) \quad (3)$$

At constant temperature of 20°C, the equation becomes (Fredlund and Rahardjo, 1993):

$$\psi = -135022 \ln(RH) \quad (4)$$

Equation (4) indicates that total suction can be determined by measuring relative humidity in the surrounding closed system. RH 100% indicates that no suction, while RH < 100% indicates the presence of suction in the system. Relative humidity measurement can be conducted by direct method such as psychrometer or by indirect method using filter paper method. The description of filter paper method will be presented in this paper, whereas psychrometer method is beyond of this study.

The principle of filter paper method in soil suction measurement is based on assumption that filter paper comes into equilibrium with soil having specific suction (Fredlund and Rahardjo, 1993). The equilibrium can be reached by transferring moisture from soil to filter paper, either through liquid flow or water vapour flow. Water, in term of liquid

or vapour may move from wet soil to dry filter paper. This movement occurs continuously until equilibrium is reached. In equilibrium, the suction of soil and filter paper will be the same. Total suction can be determined by measuring absorbed moisture by filter paper through vapour flow, whereas matric suction can be determined by measuring absorbed water through liquid flow. In short, the soil suction can be determined by measuring moisture content of filter paper.

Filter paper calibration curve should be established prior to measure soil suction. Calibration curve is a curve that relates moisture content of filter paper and suction value of the system at equilibrium. In the past, a calibration curve for particular filter paper can be use either for total or matric suction. However, recent studies indicate that for total and matric suction measurement, two separate calibration curves should be made, one for total suction and one for matric suction (Houston et al., 1994; Ridley, 1995; Bulut and Wray, 2005). In other word, total suction calibration curve will only be applicable for total suction. Likewise, matric suction calibration curve will only be applicable for measuring soil matric suction.

Referring to equation (4), calibration curve for filter paper can be made by plotting moisture content of filter papers vs. suction. Some salt solutions such as NaCl, KCl, MgNO₃, NH₄Cl, and CaSO₄ with different molality are commonly used to generate relative humidity of water vapour above solution. ASTM D 5298-94 provides relative humidity values generated from NaCl and KCl solutions at 20°C (Table 1). Molality of solution is calculated using equation (5) and (6). Figure 1 shows the linear curves of molality vs. suction which the values of R² are very close to 1.

Table 1. Relative humidity and suction values generated from NaCl and KCl at 20°C (except column 2 and 4, all data are taken from ASTM D 5298-94)

NaCl		KCl		RH	Suction	
g/1000 mg water	Molality	g/1000 mg water	Molality		KPa	Log (KPa)
1.3	0.022	1.7	0.023	0.99927	-98	1.99
3.8	0.065	5.3	0.071	0.99774	-310	2.49
13.1	0.224	17	0.228	0.99278	-980	2.99
39	0.667	52.7	0.707	0.97764	-3099	3.49
122.5	2.096	165.0	2.213	0.93008	-9800	3.99

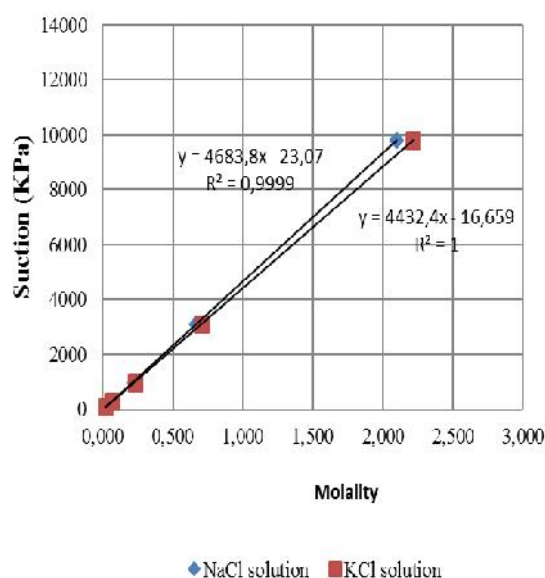


Figure 1. Suction values generated from different molality of NaCl and KCl solutions at 20°C (Data are taken from ASTM D 5298-94)

3. EXPERIMENTAL PROCEDURE

For generating suction, NaCl and KCl solution with different molality starting from 0.1 m, 0.5 m, 1 m, 2 m, 4 m, and maximum 6 m has been made. The molality of solution is calculated as:

$$\text{molality} = \frac{\text{mole of solute}}{\text{mass of solvent (kg)}} \quad (5)$$

where

$$\text{mole of solute} = \frac{\text{mass of solute}}{\text{molecular mass of solute}} \quad (6)$$

NaCl and KCl are purchased from Rowe Scientific which is having molecular mass of 58.44 and 74.55 respectively (Figure 2a). To make 1 molal NaCl solution, 1 mole NaCl is mixed with 1000 ml distilled water. 1 mole NaCl can be obtained by weighing 58.44 gr of pure NaCl. Likewise, 74.55 gr KCl is mixed with 1000 ml distilled water to make 1 molal KCl. Table 2 illustrates approximation of solute mass of NaCl and KCl for providing desired solutions. 50 ml of each solution is then poured into 130 ml glass jar (Figure 2b).

Table 2. Solute mass for different molality of NaCl and KCl

Distilled water (solvent): 1000 ml			
Solute mass (gr)		Molality	
NaCl	KCl	NaCl	KCl
6	7.5	0.103	0.101
29	37	0.496	0.496
59	75	1.010	1.006
117	149	2.002	1.999
234	298	4.004	3.997
350	447	5.989	5.996

In this study, 55 mm Whatman # 42 ashless filter paper of batch K11776722 is used. Prior to place in glass jars, filter papers are oven dried at 105°C for at least 24 hours, and placed in desiccators. Filter paper is then suspended + 10 mm above solution surface in glass jar and is held by steel wire (Figure 2c). Immediately after placing filter paper, glass jar is covered tightly by plastic lid, sealed with electric tape (Figure 2d), and put into insulated plastic box. The position of glass jars is tilted 10°-20° to avoid droplet due to condensation. The ambient temperature is controlled at near 20°C, while the temperature of insulated plastic box is checked regularly by using thermocouple thermometer. The set up of experiment is illustrated in Figure 2.

There are different equilibration periods for filter paper test. ASTM recommended using minimum 7 days for equilibration. Marinho and Oliveira (2005) suggested 7 days for 1000 – 30000 KPa, 15 days for 250 – 1000 KPa, 30 days for 100 -250 KPa, and more than 30 days for 0 – 100 KPa. However, most of studies use 7 – 14 days for equilibration period. In this study, 10 and 14 days equilibration periods are taken for NaCl and KCl solutions respectively.

After 10 days, insulated plastic box is opened and all of glass jars containing NaCl solutions are removed. Filter paper are immediately removed from glass jar, and moisture content filter paper test are conducted then by using digital balance with accuracy of 0.0001 g. Care has been taken when removing filter paper from glass jar into moisture tin to reduce moisture loss. It is performed as quickly as possible (within a few seconds) using tweezers and rubber gloves, without touching the filter paper. With the same procedure, filter paper test for KCl solution is conducted after 14 days.

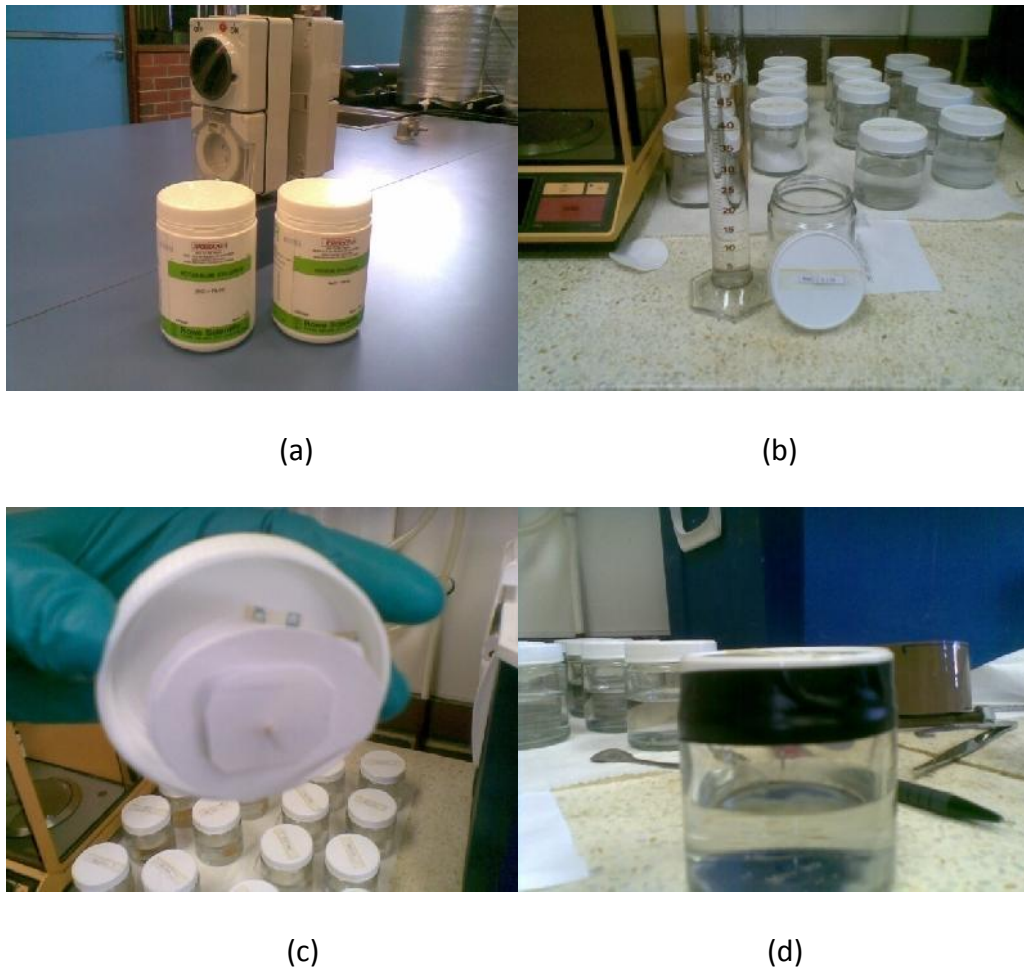


Figure 2. Photos of experimental procedure

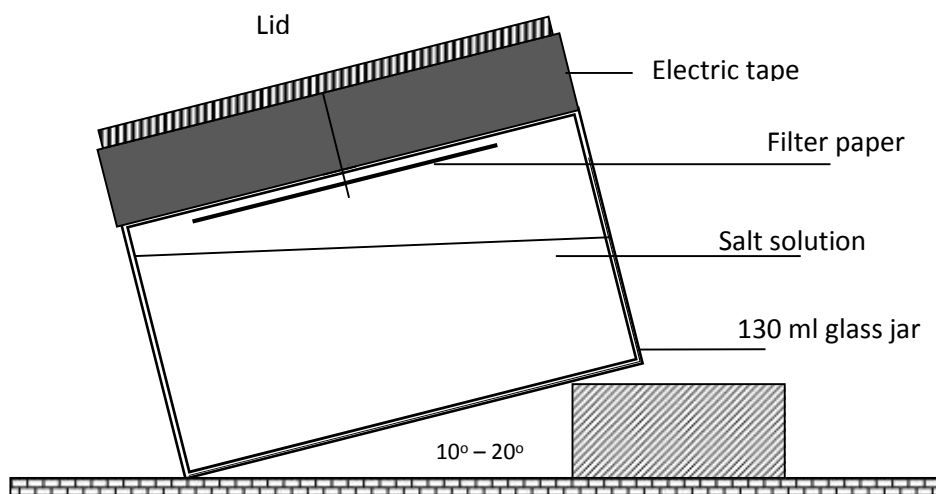


Figure 3. Experimental set up of glass jar inside insulated box

4. RESULT

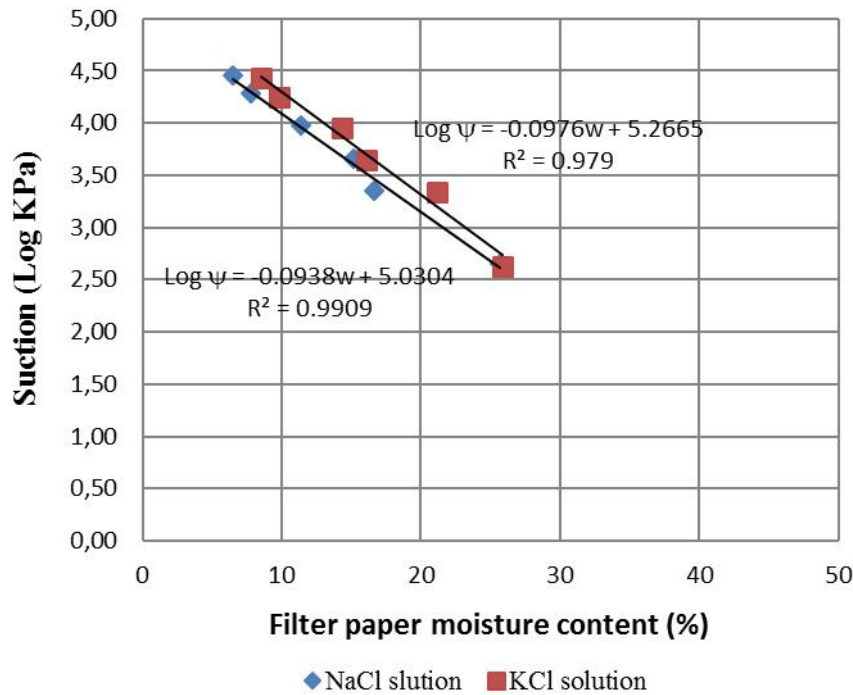
Table 3 and 4 show the calculation and result of moisture content filter paper test for total suction using NaCl and KCl solutions respectively. The suction value of each molality is taken from the curve of Figure 1. It can be observed that the higher molality of solution, the higher suction is generated. Suction can be calculated in cm water, bars, atm, pF or KPa, however, geotechnical engineers prefer to use KPa instead of other units. Figure 4 shows calibration curves where suction in Log KPa

Table 3. Moisture content filter paper test using NaCl solution (after 10 days)

Molality	m	0.1	0.5	1	2	4	6
Moisture tin no		11	15	16	20	24	29
Cold tare mass, g	Tc	38.9299	38.3518	38.3999	39.6136	40.7272	36.5816
Mass of wet FP + cold tare mass, g	M1	39.085	38.5004	38.5428	39.7538	40.8646	36.7166
Mass of dry FP + hot tare mass, g	M2	39.0472	38.4764	38.5188	39.736	40.8498	36.7065
Hot tare mass, g	Th	38.9238	38.349	38.3947	39.6102	40.7224	36.5798
Mass of dry FP, g (M2-Th)	Mf	0.1234	0.1274	0.1241	0.1258	0.1274	0.1267
Mass of water in FP, g (M1-M2-Tc+Th)	Mw	0.0317	0.0212	0.0188	0.0144	0.01	0.0083
Moisture content of FP (Mw/Mf)	W	0.2569	0.1664	0.1515	0.1145	0.0785	0.0655
Suction, kPa	ψ	454	2241	4550	9570	18939.21	28472.41
Log kPa	ψ	2.66	3.35	3.66	3.98	4.28	4.45
Suction, pF	ψ	3.66	4.35	4.66	4.98	5.28	5.45

Table 4. Moisture content filter paper test using KCl solution (after 14 days)

Molality	m	0.1	0.5	1	2	4	6
Moisture tin no		11	12	16	23	26	29
Filter paper (FP)		2	1	1	2	2	2
Cold tare mass, g	Tc	36.2799	39.7407	38.3976	37.6051	32.2350	36.5809
Mass of wet FP + cold tare mass, g	M1	36.4349	39.8940	38.5426	37.7468	32.3680	36.7142
Mass of dry FP + hot tare mass, g	M2	36.3968	39.8539	38.5190	37.7242	32.3546	36.7024
Hot tare mass, g	Th	36.2737	39.7274	38.3941	37.6003	32.2335	36.5796
Mass of dry FP, g (M2-Th)	Mf	0.1231	0.1265	0.1249	0.1239	0.1211	0.1228
Mass of water in FP, g (M1-M2-Tc+Th)	Mw	0.0319	0.0268	0.0201	0.0178	0.0119	0.0105
Moisture content of FP (Mw/Mf)	W	0.2591	0.2119	0.1609	0.1437	0.0983	0.0855
Suction, kPa	ψ	426.581	2199.541	4415.741	8848.141	17712.94	26577.74
Suction, log KPa	ψ	2.63	3.34	3.65	3.95	4.25	4.42
Suction, pF	ψ	3.63	4.34	4.65	4.95	5.25	5.42



(b)

Fig 4. Calibration Curve for Whatman #42 Filter paper using NaCl and KCl solution separately.

5. DISCUSSION

At equal concentration and equilibrium condition, NaCl generate higher suction than KCl. The different chemical properties of both salts are probably affecting the suction. It can be shown in Figure 4 that calibration curves for total suction from NaCl and KCl are a bit different. The calibration curve from KCl exhibits a bit higher compare to the one from NaCl. The curves can be expressed in equations:

$$\text{Log } \psi = -0.0938w + 5.0304 \quad \text{KPa, for NaCl} \quad (7)$$

And

$$\text{Log } \psi = -0.0976w + 5.2665 \quad \text{KPa, for KCl} \quad (8)$$

Where

w = filter paper moisture content

The difference of both curves most probably is due to the difference of equilibration period and suction source as mentioned by Leong et al. (2002). Time period of 14 days is probably better for equilibration time than 10 days. This difference is quite small ($\pm 4\%$), then it may be neglected. However, both curves have similarity in gradient. If the curves are combined, we obtain another calibration curve as shown in Figure 5. The curve can be expressed as:

$$\text{Log } \psi = -0.0933w + 5.1097 \quad \text{KPa} \quad (9)$$

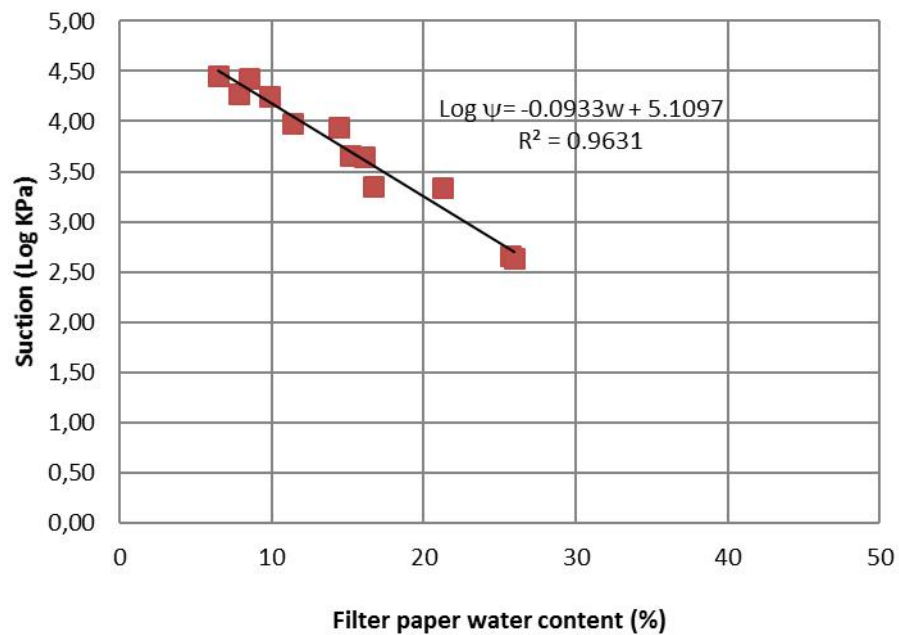


Figure 5. Calibration Curve for Whatman #42 using combination data from NaCl and KCl solutions

Calibration curve and equation (9) is very close to the curve by Leong et.al (2002) for Whatman #42 for total suction less than 26 % (equation 14). They suggested the equations for total and matric suction as:

Matric suction: $\text{Log } \psi = -0.0229w + 2.909$ $w \geq 47 \%$ (10)

$\text{Log } \psi = -0.0673w + 4.945$ $w < 47 \%$ (11)

Total suction: $\text{Log } \psi = -0.222w + 8.778$ $w \geq 26 \%$ (13)

$\text{Log } \psi = -0.0879w + 5.31$ $w < 26 \%$ (14)

Equation (9) up to equation (14) can be plotted in one are plotted in one graphic as shown in Figure 6.

In this study, the maximum water absorbed by filter paper in term of moisture content is 26 % (± 1000 KPa). Beyond this point, filter paper is not able to absorb the water anymore. The previous study conducted by several investigators also indicated that it is hard to obtain suction lower than 1000 KPa by using this method. To overcome this shortcoming, they use other apparatus such as suction plate, pressure plate, vacuum desiccators, and pressure membrane. The complete matric and total suction calibration curves may be obtained by using combination of those methods.

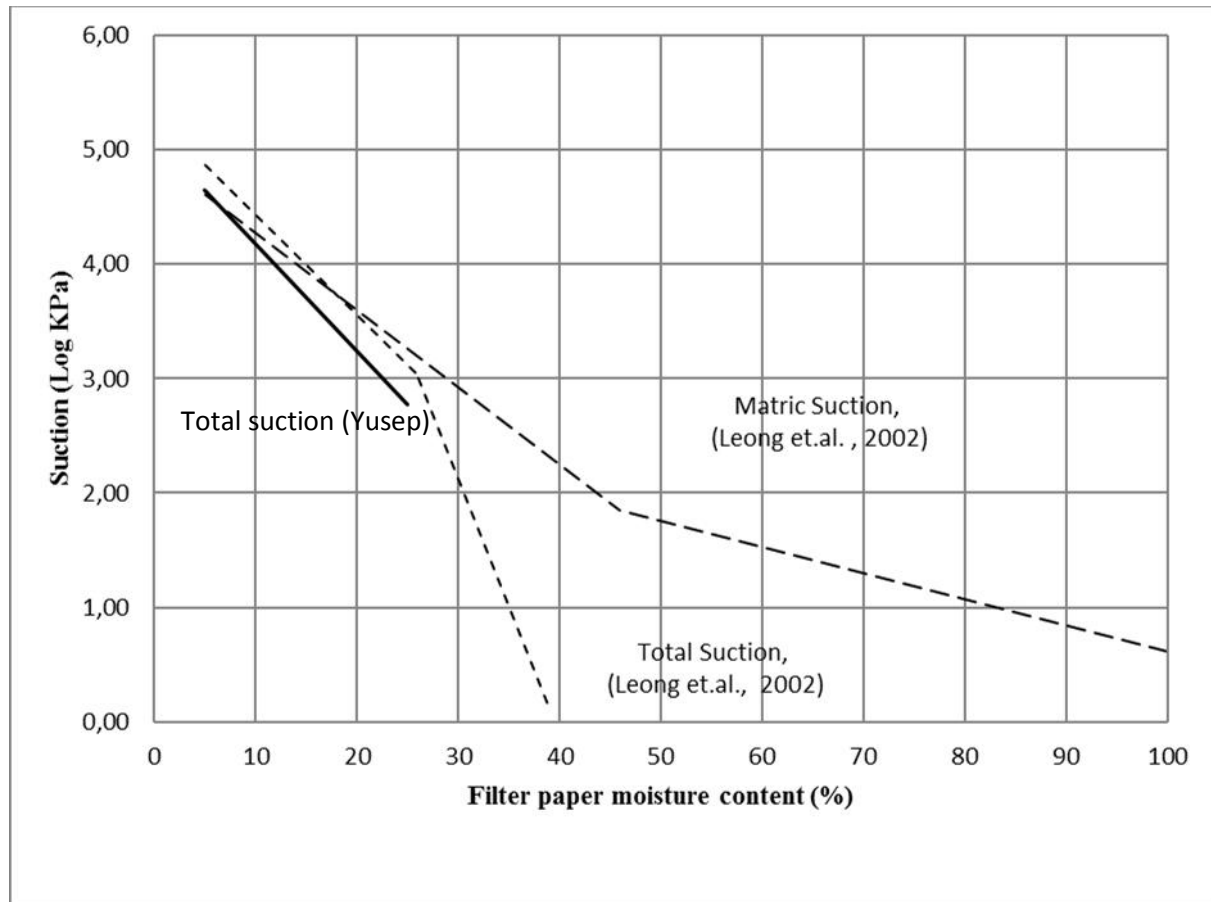


Figure 6. Calibration curve for Whatman #42. The dashed curves (----) are resulted from Leong and Rahardo (2002)

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