

Original Research Article

**PHYTOCHEMICAL AND ANTIBIOTIC TRAITS OF
(*CRASSULA OVATA*) JADE PLANT ON DIFFERENT
STRAINS OF BACTERIA**

Abstract

The *Crassula ovata* plant has been used for many years as an ornamental plant, and also as a medicinal plant in some communities like the Khoi of South Africa and in Chinese culture. Locally the plant is being used by homeowners who have it in their vicinity as a remedy for diarrhea and disinfecting wounds. However, the major problem of using this plant is its ineffectiveness to 'heal' wounds and diarrhea in most cases where it is being used. It brings so many questions in mind like does the *Crassula ovata* plants inhibit certain specific microorganisms, or is the concentration of the extract to blame, or even the method used to extract the plant. The mode of extraction used in this study involved both aqueous extraction and methanolic extraction, to ensure all plant constituents are extracted for better results. The microorganisms that were tested against the plant extracts are the major day to day sources of diarrhea and wound infection. The explants extracts are used at varying concentrations. The observable results were quantitatively analyzed to see which plant extract and at which concentration causes the most inhibition on the microorganisms. The plant extract with the most inhibition was found to be the water extraction at the concentration of $\times 10^0$, and it would be recommended that the *Crassula ovata* plant be used to the specifications as observed in the study.

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36 CHAPTER ONE

37 1.0 INTRODUCTION

38 Plants have been used as medicines throughout history. Medicinal plants are widely and
 39 successfully used on every continent. In Asia, the practice of herbal medicine is extremely well
 40 established and documented. As a result, most of the medicinal plants that have international
 41 recognition come from this region, particularly from China and India. In Europe and North
 42 America, the use of herbal medicine is increasing fast, especially for correcting imbalances
 43 caused by modern diets and lifestyles. Many people now take medicinal plant products on a daily
 44 basis, to maintain good health as much as to treat illness.

45 In Africa, attitudes towards traditional, herbal medicines vary strongly. One reason for this is the
 46 confusion between herbal medicine and witchcraft. The use of medicinal plants is sometimes
 47 associated with superstition, and therefore rejected by some people in favor of western medicine.
 48 On the other hand, there are millions of Africans who prefer traditional methods of treatment.

49 The valuable medicinal properties contained in certain plants are not in doubt. In recent years,
 50 for example, the Chinese plant *Artemisia annua*, has become the essential ingredient in a new
 51 generation of anti-malaria drugs. The plant is now being grown in East African countries to
 52 supply pharmaceutical manufacturers in Europe. The bark of the tree *Prunus africana* is used in
 53 making treatments for prostate cancer. *Sutherlandia*, a native plant of South Africa, is being
 54 increasingly recognized for its value to HIV/AIDS sufferers. Other African plants, such as
 55 Devil's Claw and African Geranium, are also gaining popularity as herbal medicines, particularly
 56 in Europe.

Medicinal plants therefore represent an important opportunity to rural communities in Africa also in Kenya, as a source of affordable medicine and as a source of income. Governments too need to be thinking about how to promote the benefits that medicinal plants have to offer, which may involve integrating herbal medicine into conventional healthcare systems. This raises important issues, such as regulation of traditional healers and ensuring certain standards are met.

Many common plants grown in Kenya have valuable medicinal properties. Paw paws, for instance, can be used to treat asthma, rheumatism and intestinal worms. Lemongrass can help in relieving fever. Sap from the *Aloe vera* is excellent for treating burns. These plants, and many others, can easily be grown in home gardens for domestic use. *Moringa oleifera* is another plant that has great potential, both in terms of home use and as a source of income. It has high levels of iron, calcium and Vitamin A, and can be used to boost the immune system, as well as treat a range of illnesses. It is normally consumed by drying the leaves and then pounding them into a powder. This can then be mixed with flours, or with other foods such as meat. For HIV/AIDS sufferers it offers an excellent source of nutrients which can help to support their immune system and slow down the advance of the disease.

Having a selection of different medicinal plant products can increase the number of customers. For example, the *Mondia whytei* processors in Kenya sell the raw roots of the plant, but also produce a powdered form. This is preferred by hospitals, which use it to promote appetite in patients and to increase milk production in nursing mothers. The powder is also used to make fortified foods for the sick. Other products for sale include mondia seeds and seedlings. Once products have been formulated for sale, gaining official recognition and approval from the authorities is valuable. In Kenya, the organization Action for Natural Medicine (NAMEDO) is working with the National Drug Authority to have its soaps, creams and oils approved. The organization is also working with the National Bureau of Standards, so that the products are standardized. This makes it much easier to market the products, for example through clinics or supermarkets.

Herbal and conventional doctors are frequently seen as rivals, having little respect for the skills and knowledge of the other. For example, hospitals and clinics can be swamped by people with relatively minor ailments, some of which might be treated with herbal remedies. This would allow hospitals to devote more of their resources to deal with serious diseases and operations.

However, for a government health ministry to promote or encourage people to use herbal medicines normally requires a radical change in thinking and policy. In The Gambia, this process is underway, with the government working to have traditional healers registered in associations. This will make it easier for their practice to be monitored, to ensure that it is in line with the national traditional medicine policy. This policy, currently in draft stage, aims to protect the rights of patients, to introduce standards for traditional medicine, and to protect the intellectual property rights of traditional healers. Integrating plant medicine into national policy involves not just the health ministry. Agriculture, environment and trade ministries will also be involved, so that farmers can be given support in growing the plants, harvesting from the wild can be controlled and quality standards introduced for those trading in medicinal plants and their products.

CHAPTER TWO

2.0 LITERATURE REVIEW

2.1 Brief description of *Crassula ovata*

2.1.1 Scientific classification and morphological description

Crassula ovata, commonly known as the jade plant or the money tree, belongs to the *Crassulaceae* or the Orpine family; they are a family of dicotyledons, a succulent plant with small pink or white flowers. They store water in their succulent leaves. (Springer 2003)

It is an evergreen plant up to 1 - 3 m tall, with thick branches and smooth, rounded, fleshy leaves that grow in opposing pairs along the branches which are also short and stubby but well-proportioned. Leaves are a rich jade green, 30 -90 mm long and 18 - 40 mm wide, egg-shaped to elliptic, often with a red margin and a somewhat pointed end. They are in opposite pairs, the one pair arranged at right angles to the next, and they are clustered towards the ends of the branches. New stem growth is the same color and texture as the leaves, but becomes brown and woody with age. Under the right conditions, they may produce small white or pink star-like flowers in

113 early spring. The flowers later develop into small capsules, each holding many tiny seeds (Gary
114 2004).

115 **2.1.2 Ecology and Distribution**

116 In *Crassula ovata*, the plant is able to maintain minimum water loss while photosynthesizing
117 efficiently through Crassulacean Acid Metabolism (CAM). The stomata are closed during the
118 day but open at night when the CO₂ taken in is stored in the form of organic crassulacean acids.
119 During the day, these acids are broken down and the CO₂ released is re-used in the
120 photosynthetic process. In this way they lose much less water yet can photosynthesize normally
121 during the daylight hours. Furthermore, during extremely dry periods they won't even open their
122 stomata at night, and will re-cycle the CO₂ within the cells. They won't be able to grow at all but
123 the cells will be kept healthy - this is known as CAM-idling (Walter *et al.*, 2012)

124 The plants succulent water-storing stems, leaves and swollen roots give it the ability to survive
125 droughts, being grazed, trampled on or knocked over, as it is able to root from any piece of stem,
126 and even a single leaf. Any discarded leaves left around the foot of the plant send down roots and
127 grow into new plants.

128 The flowers of *Crassula ovata* attract bees, wasps, flies, beetles and butterflies and also wind
129 which help disperse the fine dust-like seeds. The stems also make handy bases for wasps to build
130 their nests (Eggli 2002).

131 *Crassula ovata* is native to South Africa, and is a common houseplant all over the world, but
132 mostly occur in the Northern Hemisphere especially in dry and/or cold areas where water may be
133 scarce. *Crassula ovata* is a prominent element of the Eastern Cape and KwaZulu-Natal valley
134 thicket vegetation, together with a variety of aloes, euphorbias, *Portulacaria afra* and other
135 succulents. It occurs from Willowmore to East London and northwards to Queenstown and
136 KwaZulu-Natal where it grows on rocky hillsides (Leistner 2000).

137 In Kenya the *Crassula ovata* is found growing in areas with adequate rainfall which is well
138 distributed throughout the year. These are areas within the Central, Rift Valley, Nyanza, and few
139 areas in the Eastern region of Kenya. The *Crassula ovata* rarely grows in the North Eastern part
140 of the country due to the scarce availability of precipitation. Also in the coastal region it's very

141 rare to find this plant. There is no variation in the *Crassula ovata*'s phytochemical composition
142 regardless of where they are from. The only difference might occur in their succulence
143 depending on the geographical location which will affect water availability in the area where the
144 plant is found (Gary 2004).

145 **2.1.3 Mode of Propagation**

146 *Crassula ovata* is famously propagated either by leaf cuttings or stem cuttings. Both of these
147 types of cuttings require high humidity. In the wild, stems and leaves will often break off and fall
148 to the ground, and after a few weeks, they may grow roots and form a new plant. They can also
149 be cut and placed in a water container until roots grow usually in about two weeks, then planted
150 in soil.

151 In cultivation, new plants are made by cutting new growth (stems or leaves) and letting them dry.
152 Roots will develop in or out of soil, though inserting the stem into moist soil will increase
153 rooting. (Hudson *et al.*, 2002)

154 **2.2 Traditional Uses and Cultural Aspects**

155 Traditionally many communities have developed a habit of using the fluid extract from the
156 leaves to treat warts which are small circumscribed tumor of the outer layer of the skin. Warts
157 are flat or elevated from the surrounding skin and are firm. They are caused by forms of the
158 contagious human papilloma virus (HPV); warts vary in size and may be accompanied by pain,
159 particularly if they occur on the feet (plantar warts). The leaf of *Crassula ovata* was sliced in
160 half and attached the moist inside to the wart for a few hours, or overnight. The unsightly growth
161 would fall off with just three applications (Springer 2003).

162 In Asian cultures particularly in China (700AD), jade plant is a popular element. Medicine-men
163 prescribed a tea of the jade plant to treat symptoms of diabetes. Because of its abundance and its
164 softness in ancient times, it could easily be shaped into various forms thus it was used in the art
165 of Bonsai. The plant was spread around as luxurious gift to royalties all over the Chinese empire.

166 The jade plant is used in the practice of Feng Shui to attract the flow of money. Feng Shui is the
167 Chinese art of creating balance and harmony of energies within a space. Practitioners believe that
168 the "money tree" brings balance to the southeastern corner of a home. The jade plant is one of the

plants used in this way. A jade plant is often placed near a cash register in Chinese tradition as a way to attract prosperity (Springer 2003).

In Africa, jade leaves are boiled in milk and consumed to stop diarrhea. The Khoi and other African tribes ate the roots, they were grated and cooked after which they were eaten with thick milk. The leaves were also used medicinally, boiled in milk as a remedy for diarrhea, and used to treat epilepsy, corns and as a purgative.

In the Far East, Germany and the USA it is traditionally grown in square porcelain tubs with 'lion feet' to bring good financial luck, and has attracted more common names including the Money Tree, Penny Plant, Dollar Plant and Tree of Happiness (Doreen *et al.*, 2000).

The *Crassula ovata* plant is in Kenya mostly grown in local homesteads for its ornamental value. However some people keep this plant also for its medical values. The Kamba community believe that the juice extracted from this plant help heal burn wounds on the skin. Other communities like the Maasai use it as a relief for stomach upsets.

2.3 Diseases Controlled by *Crassula ovata* plant

Microorganisms are common inhabitants of the human skin and gut flora, soil, water, and gastrointestinal tract. However, these microorganisms can also be major causes of abnormalities in the human body system. Bacteria such as some *Staphylococcus* species live on normal skin and on mucous membranes and cause no harm. Some bacteria; however, invade normal skin, broken skin or wounds causing wound infection. The most common causative organisms associated with wound infections include *Staphylococcus aureus* and *Pseudomonas aeruginosa*.

Staphylococcus aureus is a Gram-positive bacteria and a potential pathogen. It is a leading cause of bacterial disease in humans. It can be transmitted from the nasal membranes of an asymptomatic carrier to a susceptible host. This bacterium causes Furunculosis, a medical condition in which large areas of the skin are covered in persistent boils. Folliculitis is also caused by the *Staphylococcus aureus* bacterium. It's an inflammation of one or more follicles, especially of the hair, producing small boils. These infections are commonly found in young children aged 5-12, or any immuno-suppressed person. *Staphylococcus aureus* can cause impetigo (skin infection), scalded skin syndrome and food poisoning. (Gibbons *et al.*, 1978)

Pseudomonas aeruginosa is an opportunistic pathogen of humans that can invade virtually any tissue. It is a leading cause of hospital-acquired (nosocomial) gram-negative infections, but its source is often exogenous (from outside the host). *Pseudomonas aeruginosa* causes wound infections, athlete's foot, gram negative folliculitis, chronic paronychia, and pneumonia.

(Balcht *et al.*, 1994)

Escherichia coli are a Gram-negative, rod-shaped bacterium, a consistent resident of the small intestine. Some strains of *Escherichia coli* are pathogens that cause intestinal infections, urinary tract infections and neonatal meningitis. Some serotypes can cause serious food poisoning in humans. The harmless strains are part of the normal flora of the gut, and can benefit their hosts by producing vitamin K₂, and by preventing the establishment of pathogenic bacteria within the intestine. *Escherichia coli* and related bacteria constitute about 0.1% of gut flora, and fecal–oral transmission is the major route through which pathogenic strains of the bacterium cause disease.

(Vogt and Dippold 2005)

Candidiasis is an overgrowth of a fungus hence causing irritation and swelling. Pathogenicity among yeast-like fungus is extremely variable; however, the most virulent is *Candida albicans*. Involvement of the *Candida albicans* may be localized to the mouth, throat, skin, scalp, vagina, fingers, toes, nails, bronchi, lungs or gastrointestinal tract. It may also be systemic as in septicemia (circulating in the blood and causing damage to blood vessels and sometimes blood cells), endocarditis and meningitis. Factors predisposing people to candidiasis include AIDS, burn patients, young individual, pregnancy, oral birth control, high fruit diets, steroids, antibiotic therapy, immunosuppressants, cancer treatments, heart surgery, genetic deficiency, endocrine deficiency diabetes, use of catheters, and use of dirty needles.

Bacillus subtilis cells are rod-shaped, Gram-positive bacteria that are naturally found in soil and vegetation. *Bacillus subtilis* bacteria are non-pathogenic. They can contaminate food; however, they seldom result in food poisoning (Balcht *et al.*, 1994).

2.4 Statement of the problem

The main problem or general question pertaining the *Crassula ovata* is whether it has relevantly effective antibiotic or antifungal traits. *Crassula ovata* plants are mostly used as house plants and

do not have many commercial uses other than for ornamental value. However it's usually integrated into most homesteads also because of its healing properties or medicinal values. In many occurrences, the plant extracts do not always treat some stomach upsets or even fresh wounds, despite its prominent successful use in other past communities from different nations. Even after continuous application of the plant extracts, the stomach upset or wound still continues to persist.

The question left unanswered is whether the extraction procedure is efficient, or maybe the concentration of the extract is too high or too low to be effective. Or perhaps which part of the *Crassula ovata* plant is most effective to use.

The *Crassula ovata* has had successful ratings in its past ancient uses, but very much limited success in present time. There is need to research further why this has come to be. Among the many reasons for these changes might be the fact that since these plants were originally from what we now know as highly productive nations (China and South Africa), due to the industrial advancements and resulting increased environmental pollution, the plant genome has been altered. This could to a great deal alter the overall efficiency of the plants antibiotic traits.

2.5 Justification

This research was conducted so as to test the medicinal value of *Crassula ovata* plant, and whether it has any phytochemical components which inhibit growth of microorganisms. There is also the need for more research on the evolution of the specific microorganisms *Crassula ovata* is said to inhibit. This is because it is a known fact that these microorganisms and others are mutating almost every day, thereby increasing their survival and reducing the effects of any antibiotic stimuli. Unfortunately, no much research has been put in effect to solve this riddle.

2.6 Objectives

2.6.1 Broad objectives

- To determine the phytochemical components and antibiotic traits of the *Crassula ovata* plant.

252 2.6.2 Specific objectives

- 253 1. To determine the phytochemical components of the *Crassula ovata* plant.
- 254 2. To test the antibiotic activity of the *Crassula ovata* plant extracts against a range of
- 255 selected microorganisms.

256 2.7 Hypothesis

257 1.7.1 Null hypothesis

258 There is no difference in the phytochemical components of the *Crassula ovata* plant and the

259 proliferation of the selected microorganisms.

260 CHAPTER THREE

261 3.0 MATERIALS AND METHODS

262 3.1 Sample and Sampling Technique

263 The samples for experimentation include the leaves and stem of the *Crassula ovata* plant. The

264 *Crassula ovata* plant species were acquired randomly from a local homestead in Ruiru, Kenya.

265 3.2 Extraction of crude extracts from the plant

266 This was carried out according to Walter *et al.*, (2012).

267 3.2.1 Aqueous extraction

268 2 grams of the plants leaves and stem were obtained and crushed using a pestle and mortar while

269 adding 100ml sterile distilled water to dissolve the crude extracts. The extracts were then put into

270 sterile conical flasks and stored at room temperature. Serial dilution of the extract was then done

271 four to five times starting with a concentration of $\times 10^0$ to $\times 10^{-4}$.

272 3.2.2 Methanolic extraction

273 2 grams of the plants leaves and stem were obtained and cut into smaller pieces and put in 100ml

274 of 90% methanol to dissolve the crude extracts and left overnight. The plant material was then

275 separated from the methanol by washing with 100ml of sterile distilled water. Serial dilution of

276 the extract was then done four to five times starting with a concentration of $\times 10^0$ to $\times 10^{-4}$.

277 Small circular paper discs were put into the containers containing the different plant extracts
278 from the water extraction and the methanolic extraction. The discs with the methanolic extracts
279 were then put in an oven at 40°C for 30minutes.

280 **3.3 Phytochemical Tests**

281 The phytochemical tests were carried out as per procedures by Seidel (2012).

282 **3.3.1 Alkaloid Test**

283 0.05g of the sample was added to 1%HCL and filtered. The filtrate is tested carefully with
284 various alkaloid reagents as follows;

285 **3.3.1.1 Mayer's test**

286 To 1ml of the filtrate, a drop or two of Mayer's reagent was added by the side of the test tube. A
287 white or creamy precipitate indicates the test as positive.

288 **3.3.1.2 Dragendorff's test**

289 To 1ml of the filtrate 1 or 2mls of Dragendorff's reagent was added. A prominent yellow
290 precipitate confirms the test as positive.

291 **3.3.2 Carbohydrate Test**

292 **3.3.2.1 Barfoed's test**

293 To 1ml of filtrate, 1ml of Barfoed's reagent was added and heated in a boiling water bath for 2
294 minutes. A red precipitate confirms sugar presence.

295 **3.3.2.2 Benedict's test**

296 To 0.5ml of filtrate, 0.5ml Benedict's reagent was added and the mixture heated in a boiling
297 water bath for 2 minutes. A characteristic colored precipitate confirms the presence of sugar.

298 **3.3.3 Detection of Saponins**

299 1ml of plant extracts were dissolved in anhydride-tetrachloride to which 4 drops of concentrated
300 sulfuric acid was added to the mixture. A blue, green or red color accompanied by a pink ring
301 shows presence of Saponins.

3.3.4 Flavanoids Test

1ml of the extract was put into a test tube followed by addition of Hydrochloric acid (4 drops) and Magnesium turnings. Development of a pink or magenta red indicates the presense of Flavanoids.

3.3.5 Tannins Test

1ml of the crude extract was dissolved in water which contains 1% gelatin and 10% NaCl. The presence of tannins is indicated by the presence of a blackish blue color. Catecol tannins are indicated by a greenish black coloration.

3.3.6 Sterols and Steroids Test

1ml of the extract was put in a test tube in which 0.5ml sulfuric acid, acetic anhydride and chloroform in similar amounts were added. A red coloration would indicate presence of sterols. A green color indicates presence of steroids.

3.4 Media preparation and incorporation with bacteria

Based on Baker et al., (2001), Mueller Hinton agar was prepared by measuring 28.5g and dissolving it in 750ml distilled water. Nutrient broth was also prepared and put into glass bottles. The prepared media, nutrient broth, pipette tips, paper discs, distilled water, the pestle and mortar were autoclaved at 121°C for 15 minutes. The agar was left to cool to about 40-37°C then aseptically poured into sterile Petri dishes. This was done on the bench, using flame to keep media bottle sterile.

Sterile nutrient broth was inoculated with fresh bacteria strains. Bacteria was picked from a frozen culture by scratching the sterile loop across the surface of the culture or they were picked from a liquid culture by immersing loop in it. The bacteria were evenly spread across the surface of the plate using a glass spreader.

3.4.1 Disc diffusion method

Several circular sterile paper discs, were each infused with the different concentrations of the crude extracts, then evenly spaced over the surface of the plate. The discs were gently pushed down into the agar to make contact with the bacteria. The plates were left to grow overnight in an incubator at 37°C. Colonies would be visible after 12-16 hours growth at 37°C. Plates should be inverted in the incubator to prevent condensation from dripping on the colonies.

331 The colonies that form would be then counted as colonies per unit after incubation. The zones of
 332 inhibition were also measured on each plate. The minimal inhibitory concentration (MIC) of the
 333 crude extract to specific bacteria can then be determined.

334 CHAPTER FOUR

335 4.0 RESULTS AND DISCUSSION

336 4.1 PHYTOCHEMICAL TESTS RESULTS

337 The biologically active compounds of the *Crassula ovata* plant are tested so as to draw valuable
 338 conclusions from the observed results.

339 **Table 1: Phytochemical test results of the *Crassula ovata* plant crude extracts**

TEST	OBSERVATIONS	OBSERVATIONS	
	Aqueous extracts	Methanolic extracts	
Alkaloid test			
i. Mayer's test	+	-	
ii. Dragendorff's test	-	-	
Flavanoids test	-	-	
Sterols and steroids Test	+	+	
Saponins Test	+	+	
Tannins Test	-	-	
Carbohydrate Test			
i. Barfoed's Test	+	+	
ii. Benedict's Test			

	+	+
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340 The phytochemical screening of *Crassula ovata* stem and root extracts revealed the presence of
 341 saponins, steroids, Carbohydrates and alkaloids.

342 4.2 ANTIMICROBIAL ACTIVITY RESULTS

343 **Table 2 Observations for the aqueous plant leaf extracts at various concentrations**

BACTERIA	Inhibition in mm per dilution				
	$\times 10^0$	$\times 10^{-1}$	$\times 10^{-2}$	$\times 10^{-3}$	$\times 10^{-4}$
<i>Bacillus subtilis</i>	0	0	0	0	0
<i>Escherichia coli</i>	6.4	6.2	6.1	6.1	6.1
<i>Candida albicans</i>	0	0	0	0	0
<i>Staphylococcus aureus</i>	0	0	0	0	0--
<i>Pseudomonas aeruginosa</i>	0	0	0	0	0

344 The zones of inhibition of the different microorganisms after exposure to aqueous plant leaf
 345 extracts. The *E coli* bacteria showed the only inhibition with the highest at the normal
 346 concentration of $\times 10^0$.

347 **Table 3 Observations for the aqueous plant stem extracts at various concentrations**

BACTERIA	Inhibition in mm per dilution
----------	-------------------------------

	$\times 10^0$	$\times 10^{-1}$	$\times 10^{-2}$	$\times 10^{-3}$	$\times 10^{-4}$
<i>Bacillus subtilis</i>	0	0	0	0	0
<i>Escherichia coli</i>	6.1	0	0	0	0
<i>Candida albicans</i>	0	0	0	0	0
<i>Staphylococcus aureus</i>	0	0	0	0	0
<i>Pseudomonas aeruginosa</i>	0	0	0	0	0

348 The zones of inhibition of the different microorganisms after exposure to aqueous plant stem
349 extracts. The *E coli* bacteria showed the only inhibition at the normal concentration of $\times 10^0$.

350 **Table 4 Observations for the methanolic plant leaf extracts at various concentrations**

BACTERIA	Inhibition in mm per dilution				
	$\times 10^0$	$\times 10^{-1}$	$\times 10^{-2}$	$\times 10^{-3}$	$\times 10^{-4}$
<i>Bacillus subtilis</i>	0	0	0	0	0
<i>Escherichia coli</i>	6.5	6.2	0	0	0
<i>Candida</i>	0	0	0	0	0

<i>albicans</i>					
<i>Staphylococcus aureus</i>	0	0	0	0	0
<i>Pseudomonas aeruginosa</i>	0	0	0	0	0

351 The zones of inhibition of the different microorganisms after exposure to methanolic plant leaf
 352 extracts. The *E coli* bacteria showed the only inhibition with the highest at the normal
 353 concentration of $\times 10^0$.

354 **Table 5 Observations for the methanolic plant stem extracts at various concentrations**

BACTERIA	Inhibition in mm per dilution				
	$\times 10^0$	$\times 10^{-1}$	$\times 10^{-2}$	$\times 10^{-3}$	$\times 10^{-4}$
<i>Bacillus subtilis</i>	0	0	0	0	0
<i>Escherichia coli</i>	6.1	0	0	0	0
<i>Candida albicans</i>	0	0	0	0	0
<i>Staphylococcus aureus</i>	0	0	0	0	0
<i>Pseudomonas aeruginosa</i>	0	0	0	0	0

355 The zones of inhibition of the different microorganisms after exposure to methanolic plant stem
 356 extracts. The *E coli* bacteria showed the only inhibition at the normal concentration of $\times 10^0$.

357 **Table 6 Anova (Single Factor) analysis of methanolic and aqueous extracts of *Crassula***
 358 ***ovata* plant**

SUMMARY

<i>Groups</i>	<i>Count</i>	<i>Sum</i>	<i>Average</i>	<i>Variance</i>
Escherichia coli	2	12.3	6.15	0.005
Bacillus subtilis	2	0	0	0
Candida albicans	2	0	0	0
Staphylococcus aureus	2	0	0	0
Pseudomonas aeruginosa	2	0	0	0

ANOVA

<i>Source of Variation</i>	<i>SS</i>	<i>df</i>	<i>MS</i>	<i>F</i>	<i>P-</i> <i>value</i>	<i>F crit</i>
Between Groups	60.516	4	15.129	15129	2.17E-10	5.192168
Within Groups	0.005	5	0.001			
Total	60.521	9				

359 The anova analysis of the effects of the plant extract on the various microorganisms. These
 360 shows a difference in the calculated F value and tabulated F value.

361

362 **4.3 DISCUSSION**

363 Based on the observations made during the study, it was observed that the *Crassula ovata* plant
 364 had active components of alkaloids, carbohydrates, sterols, steroids and saponins. These are
 365 active chemical components that are involved in inhibition of microbial activity. However the
 366 degree of the effect of these active components depends on the plant species and the overall
 367 concentration used.

368 The antimicrobial activity of the *Crassula ovata* leaf and stem extracts were studied at different
 369 concentrations against four pathogenic bacterial strains and one fungal strain. The antimicrobial

activities of the extracts increased linearly with increase in concentration of the extracts. The *Escherichia coli* bacteria were more sensitive than any other microbe to the plants extracts especially the leaf aqueous extracts. The growth inhibition zones measured an average of 6.2 mm for all the sensitive microbes. The results show that the aqueous leaf extracts of *Crassula ovata* were found to be more effective against *Escherichia coli*, but no effectiveness on the other microbes tested.

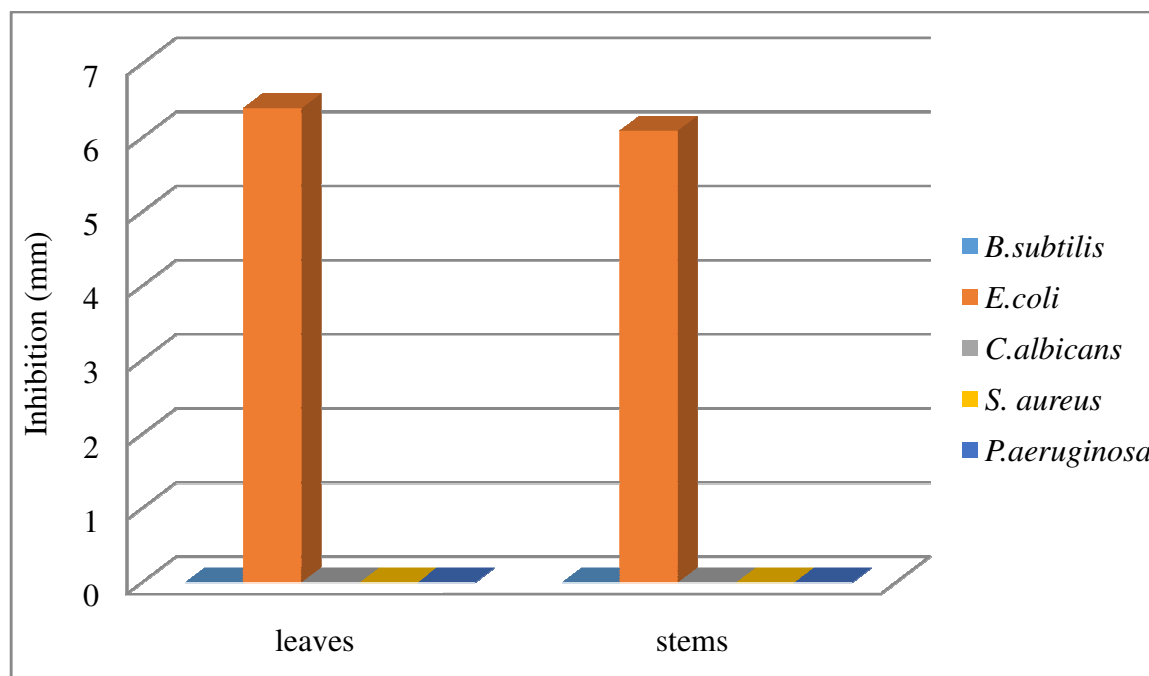


Figure 1 Antimicrobial activity against different microbes exposed to methanolic extracts
The methanolic extracts only showed inhibition of *E. coli* bacteria at concentration $\times 10^0$ only. No inhibition was observed for the other microorganisms.

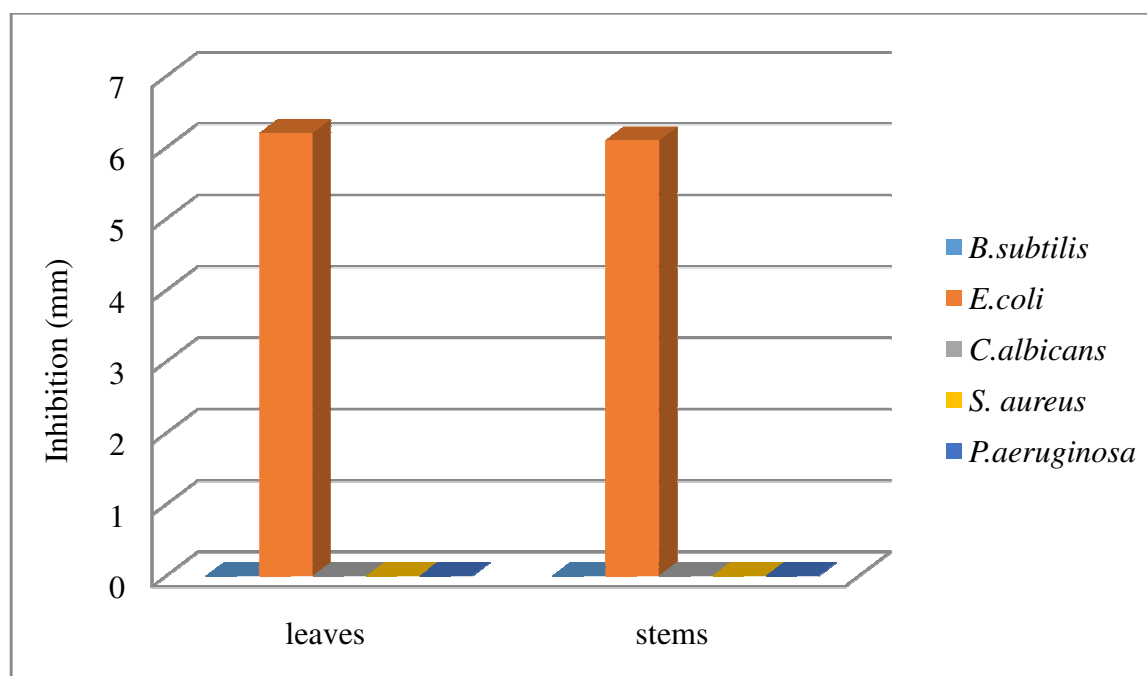


Figure 2 Antimicrobial activity against different microbes exposed to aqueous extracts
The aqueous extracts showed an average inhibition on only *E. coli* bacteria alone.

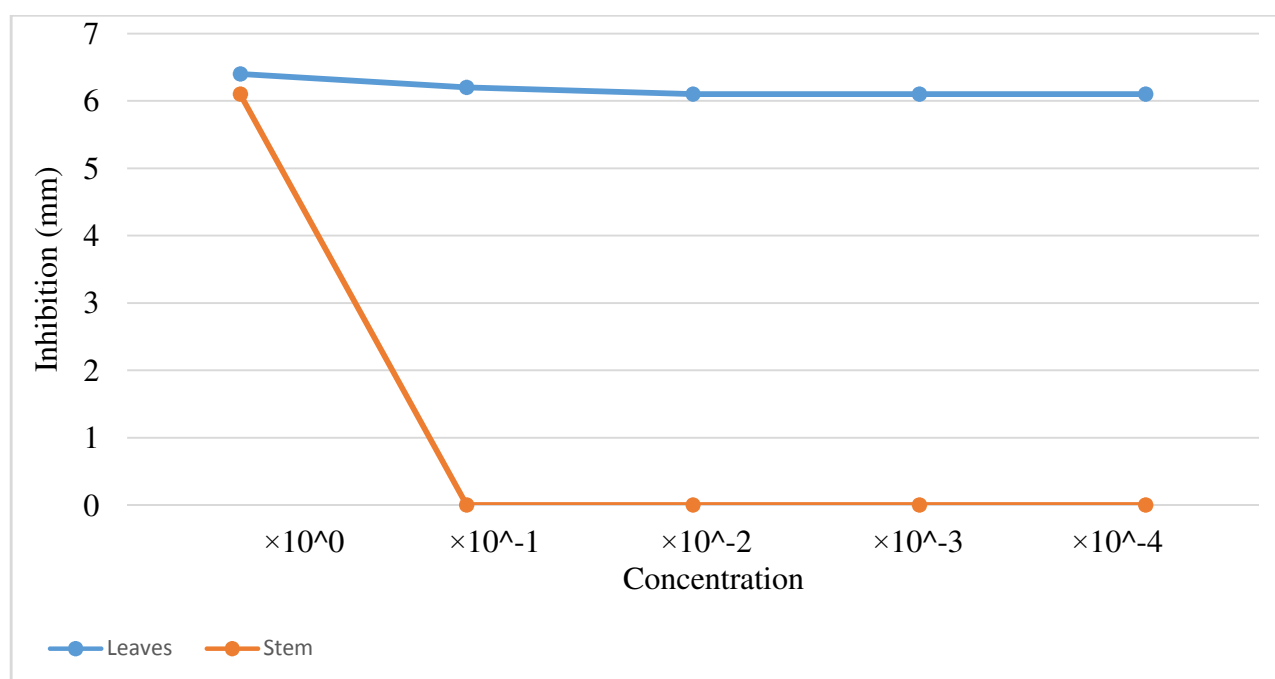


Figure 3: A Chart of Inhibition against Concentration of *E. coli* by aqueous plant extracts
The bacteria *E. coli* was greatly inhibited at $\times 10^0$ concentration, with leaf extract being at 6.4 mm and stem extracts at 6.2 mm. Inhibition declined for the stem extracts instantaneously while for the leaf extracts it was gradual up to 6.1 mm.

388 The aqueous extracts of *Crassula ovata* showed strong activity against *Escherichia coli*. The
389 results also revealed the presence of different phytochemical compounds with biological activity
390 that can be of valuable therapeutic index. It has been shown from earlier experiments that plants
391 rich in phenolic compounds have been shown to have antimicrobial activities.

392 From the anova analysis, the tabulated F value was lesser than the calculated F values did not
393 match, and hence the null hypothesis had to be rejected. The aqueous leaf extracts of the
394 *Crassula ovata* plant gave a promising effectiveness of antimicrobial growth on *Escherichia coli*
395 bacteria alone. This is attributed to the presence of alkaloids like Berberine and Sanguinarine,
396 and saponins as observed during the phytochemical testing of the plant. This showed that the
397 *Crassula ovata* plant is effective only to the *Escherichia coli* bacteria. In this light, more
398 bioprospecting questions arise on whether the plant extracts can be manipulated further to
399 completely inhibit *Escherichia coli* growth and development.

400 The *Crassula ovata* plant is a common home plant in most parts of Kenya. However, the claims
401 that it can heal wounds are most likely not true. Scientific experimentation carried out in this
402 study helps prove that point. However, due to the plants effect on *Escherichia coli*, the *Crassula*
403 *ovata* plant's potential to control stomach upset is yet to be further looked into.

404 CHAPTER FIVE

405 5.0 CONCLUSIONS AND RECOMMENDATIONS

406 5.1 CONCLUSIONS

407 It is evident from the results that the plant extracts of the *Crassula ovata* plant, both from the
408 leaves and stems, methanolic and aqueous, are only able to inhibit the *Escherichia coli* bacteria.
409 Also being a gram negative bacteria, *Pseudomonas aeruginosa* was not affected by the plant
410 extracts. Meaning that there was an active compound in the plants extracts that acted specifically
411 against *E. coli* bacteria. This results also led to the rejection of the null hypothesis since a
412 significant difference was observed in the microbial proliferation and the active compounds in
413 the *Crassula ovata* plant. Other than *Escherichia coli*, the *Crassula ovata* plant is none effective
414 to the other microbes that were tested against.

The objectives of this study were met, both the broad and specific objectives. There were active phytochemical compounds in the *Crassula ovata* plant. This included the saponins, steroids, Carbohydrates and alkaloids. The plant was found to have an antimicrobial effect on *Escherichia coli* bacteria.

This study proved that there are active phytochemical compounds in the *Crassula ovata* plant, and that these compounds have a relatively minimal effect on microbial activity.

5.2 RECOMMENDATIONS

Since the *Crassula ovata* plant is effective to inhibit growth of only *Escherichia coli* bacteria. It would be highly recommended that further research is done to ascertain to which degree the *Crassula ovata* plant extracts can inhibit the *Escherichia coli* bacteria.

Other recommendations that the author would suggest include;

- Research on different *Crassula* plant species varieties from different locations and how effective they inhibit different microorganisms.
- Use of more gram negative bacteria against the *Crassula ovata* extracts.
- Isolating the *Crassula ovata*'s active compound inhibiting the *E.coli* bacteria and molecularly engineer it using bioinformatics tools to test its potential as a possible drug.

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