

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.

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

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

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

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

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

98

99

CHAPTER TWO

100

2.0 LITERATURE REVIEW

101 **2.1 Brief description of *Crassula ovata***

102 **2.1.1 Scientific classification and morphological description**

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

106 It is an evergreen plant up to 1 - 3 m tall, with thick branches and smooth, rounded, fleshy leaves
107 that grow in opposing pairs along the branches which are also short and stubby but well-
108 proportioned. Leaves are a rich jade green, 30 -90 mm long and 18 - 40 mm wide, egg-shaped to
109 elliptic, often with a red margin and a somewhat pointed end. They are in opposite pairs, the one
110 pair arranged at right angles to the next, and they are clustered towards the ends of the branches.
111 New stem growth is the same color and texture as the leaves, but becomes brown and woody
112 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

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

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

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

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

182 **2.3 Diseases Controlled by *Crassula ovata* plant**

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

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

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

201 (Balcht *et al.*, 1994)

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

209 (Vogt and Dippold 2005)

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

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

222 **2.4 Statement of the problem**

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

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

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

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

240 **2.5 Justification**

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

247

248 **2.6 Objectives**

249 **2.6.1 Broad objectives**

- 250 • To determine the phytochemical components and antibiotic traits of the *Crassula ovata*
251 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.

302 3.3.4 Flavanoids Test

303 1ml of the extract was put into a test tube followed by addition of Hydrochloric acid (4 drops)
304 and Magnesium turnings. Development of a pink or magenta red indicates the presence of
305 Flavanoids.

306 3.3.5 Tannins Test

307 1ml of the crude extract was dissolved in water which contains 1% gelatin and 10% NaCl. The
308 presence of tannins is indicated by the presence of a blackish blue color. Catecol tannins are
309 indicated by a greenish black coloration.

310 3.3.6 Sterols and Steroids Test

311 1ml of the extract was put in a test tube in which 0.5ml sulfuric acid, acetic anhydride and
312 chloroform in similar amounts were added. A red coloration would indicate presence of sterols.
313 A green color indicates presence of steroids.

314 3.4 Media preparation and incorporation with bacteria

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

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

325 3.4.1 Disc diffusion method

326 Several circular sterile paper discs, were each infused with the different concentrations of the
327 crude extracts, then evenly spaced over the surface of the plate. The discs were gently pushed
328 down into the agar to make contact with the bacteria. The plates were left to grow overnight in
329 an incubator at 37°C. Colonies would be visible after 12-16 hours growth at 37°C. Plates should
330 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>
<i>Escherichia coli</i>	2	12.3	6.15	0.005
<i>Bacillus subtilis</i>	2	0	0	0
<i>Candida albicans</i>	2	0	0	0
<i>Staphylococcus aureus</i>	2	0	0	0
<i>Pseudomonas aeruginosa</i>	2	0	0	0

ANOVA

<i>Source of Variation</i>	<i>SS</i>	<i>df</i>	<i>MS</i>	<i>F</i>	<i>P-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.

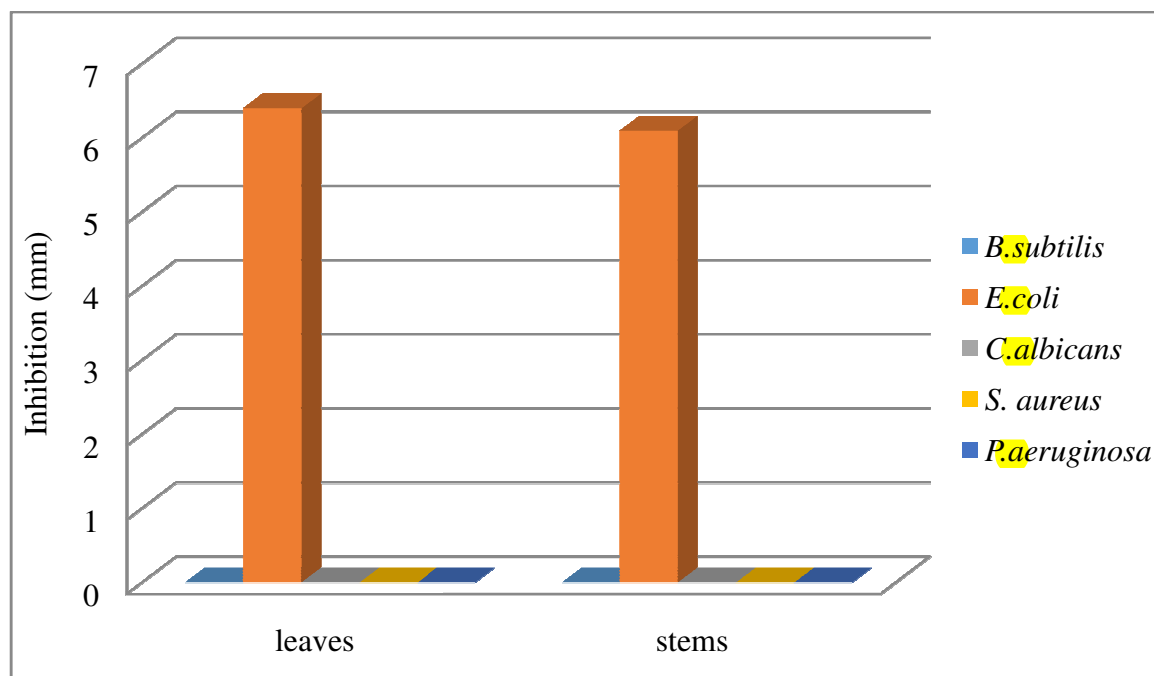
361

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

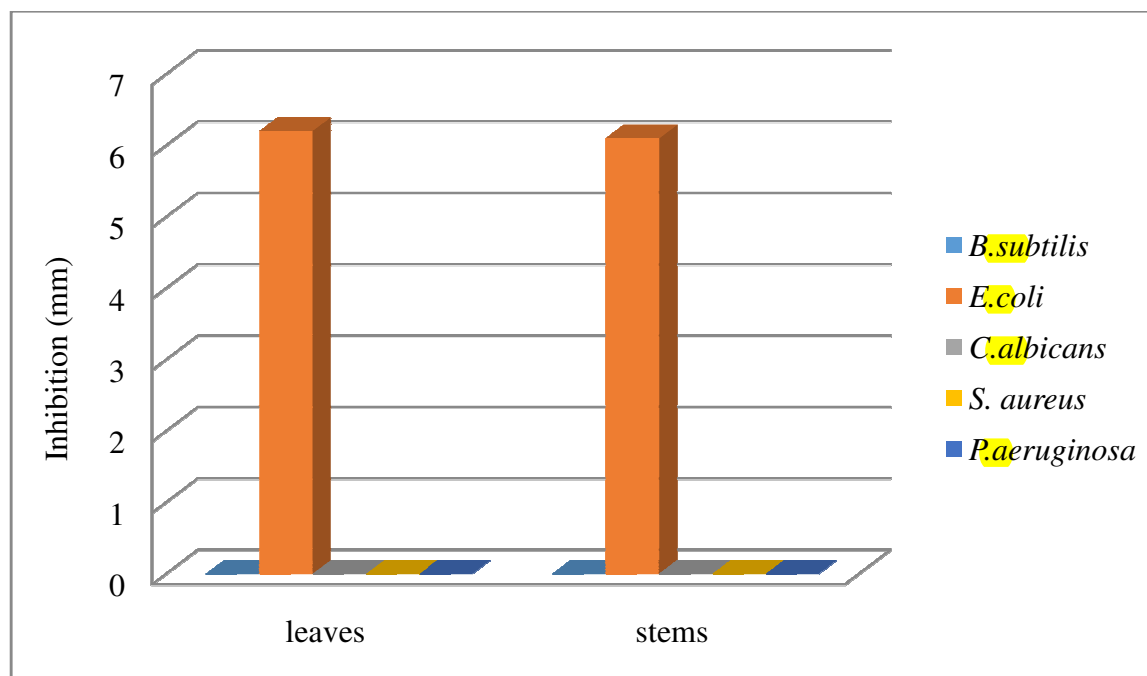
370 activities of the extracts increased linearly with increase in concentration of the extracts. The
 371 *Escherichia coli* bacteria were more sensitive than any other microbe to the plants extracts
 372 especially the leaf aqueous extracts. The growth inhibition zones measured an average of 6.2 mm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



376

377 **Figure 1 Antimicrobial activity against different microbes exposed to methanolic extracts**

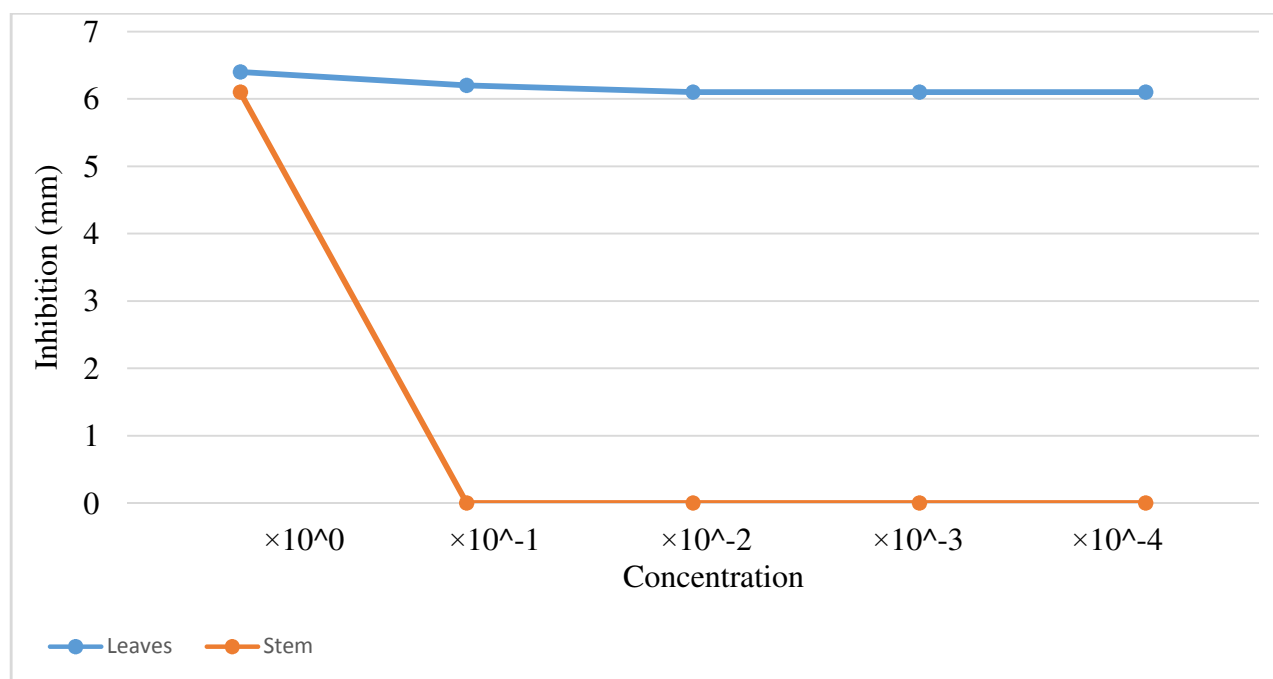
378 The methanolic extracts only showed inhibition of *E. coli* bacteria at concentration $\times 10^0$ only.
 379 No inhibition was observed for the other microorganisms.



380

381 **Figure 2 Antimicrobial activity against different microbes exposed to aqueous extracts**

382 The aqueous extracts showed an average inhibition on only *E. coli* bacteria alone.



383

384 **Figure 3: A Chart of Inhibition against Concentration of *E. coli* by aqueous plant extracts**

385 The bacteria *E. coli* was greatly inhibited at $\times 10^0$ concentration, with leaf extract being at 6.4
 386 mm and stem extracts at 6.2 mm. Inhibition declined for the stem extracts instantaneously while
 387 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

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.

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

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

421 5.2 RECOMMENDATIONS

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

425 Other recommendations that the author would suggest include;

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

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