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# A Review on Medicinal Plants Against Some Human Pathogenic Bacteria (E. coli, S.dysentery, S.typhi, P.aeruginosa and S. aureus) in South East Ethiopia

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# **INTRODUCTION**

# 1.1. Historical use of medicinal plants as medicine

In many parts of the world, medicinal plants have been used as traditional treatments for various human ailments for thousands of years. The use of plant and its products has a long history in maintaining human health that began with folk medicine and through the years has been incorporated into traditional and allopathic medicine (Gislene et al., 2000; Dubey et al., 2011). Plants have a long history of use in treatment and management of different diseases all over the world since ancient times and about 25% of current drugs are derived from plants (Wanyoike et al., 2004). In certain African countries up to 90% of the population relies exclusively on plants as sources of medicines (Hostetman et al., 2000).

Medicinal plants have been recognized as potential sources of new compounds for therapeutic use. Findings from researchers and pharmaceutical entrepreneurs have pointed out that ethno botanically derived compounds have greater activity than compounds derived from random screening and thus a greater potential for novel products developed (Njoroge and Bussmann, 2006). Natural products as pure compounds or standardized plant extracts provide an unlimited opportunities for new drug leads due to their unmatched availability of chemical diversity (Parekh and Chanda, 2007). Natural products have been used in traditional medicine all over the world for thousands of years and they predate the introduction of antibiotics and other modern drugs (Balunas and Kinghorn, 2005, Parekh and Chanda, 2007). Plants are rich in a wide variety of secondary metabolites called phytochemicals such as tannins, alkaloids, and flavonoids that have been found to have antimicrobial properties. For example, the essential oil and eugenol purified from Ocimum gratissimum has been reported to treat pneumonia, diarrhea and conjunctivitis (Nakamura et al., 1999). These evidences contribute to support and quantify the importance of screening natural products.

# 1.2. Contributions of plant extract compounds in the field of medicine

Plants are known to produce a variety of compounds to protect themselves against a variety of pathogens. It is expected that plant extracts showing target sites other than those used by antibiotics will be active against drug resistant pathogens. These types of plants that are used for treatment of different diseases are so called medicinal plants (Ahmad and Beg, 2001). Medicinal plants represent a rich source of antimicrobial agents. Plants are used medicinally in different countries and are a source of many potent and powerful drugs (Srivastava *et al.*, 1996). A wide range of medicinal plant parts is used for extract as raw drugs and they possess varied medicinal properties. The different parts used include root, stem, flower, fruit, twigs exudates and modified plant organs.

Natural products of higher plants are the probable sources of antimicrobial agents which have added advantages of being safe and biodegradable (Adenisa *et al.*, 2000). Natural products as pure compounds or standardized plant extracts provide an unlimited opportunities for new drug leads due to their unmatched availability of chemical diversity (Parekh and Chanda, 2007). Many plants species have been used because of their antimicrobial traits ; pharmacological properties as they are known to posses various secondary metabolites called phytochemicals like glycosides, saponins, flavonoids, steroids, tannins, alkaloids, tirpenes which is therefore, should be utilized to combat the disease causing pathogens (Lalitha *et al.*, 2010; Hussain *et al.* 2011).

# 1.3. Significance of extraction of medicinally valuable compounds from plants

With the advancement in Science and Technology, remarkable progress has been made in the field of medicine with the discoveries of many natural and synthetic drugs (Preethi *et al.*, 2010). Plants may serve as natural blue prints in the development of new drugs or as phytomedicines to be used to treat disease (Abubakar *et al.*, 2008). Antibiotics are undeniably one of the most important therapeutic discoveries of the 20<sup>th</sup> century that had effectiveness against serious bacterial infections. However, only one third of the infectious diseases known have been treated from these synthetic products (Sharma, 2011).

Although hundreds of plant species have been tested for antimicrobial properties, the vast majority of have not been adequately evaluated. Consequently, in recent time researchers have paid attention to advanced phytomedicines and biologically active compounds isolated from plant species used in herbal medicines with acceptable therapeutic index for the development of novel drugs (Pavithra *et al.*, 2010).

# 1.4. The in vitro antibacterial activity assay

Introduction and development of several new and highly specific in vitro bioassay techniques, chromatographic methods, spectroscopic techniques and other standardized pharmacological methods have made much easier to screen, isolate and identify potential drug compounds quickly and precisely from natural sources to alleviate human illnesses (Satyajit et al., 2001). Antimicrobial susceptibility tests measure the ability of an antimicrobial agent to inhibit bacterial growth in vitro. This ability may be estimated by either the dilution method or the diffusion method (WHO, 2003). In vitro antimicrobial test is preferable method to carry out experiments using the crude powder of each plant extract by plate diffusion method (Ieven et al., 1979). Plants are important source of potentially useful structures for the development of new chemothera peutic agents. The first step towards this goal is the in vitro antibacterial activity assay (Tona, 1998; Mahesh and Satish, 2008).

There are many methods to test the effectiveness of antimicrobial chemicals, including the agar well diffusion, disk diffusion and dilution assay. In disc diffusion assay, small filter disks are impregnated with the chemical to be tested, and are placed on a plate inoculated to form a bacterial lawn (even, confluent bacterial growth). The plates are incubated to allow growth of the bacteria and time for the chemicals to diffuse into the agar. As a chemical diffuses into the agar, it becomes less concentrated. If an organism is susceptible to a chemical, a clear zone of inhibition will appear around the disk where the growth has been inhibited. The size of this zone of inhibition depends on the sensitivity of the bacteria to the specific chemical and the chemical's ability to diffuse through the agar (Cain et al., 2013). Disc-diffusion susceptibility tests may give unreliable results, if the appropriate technique is not strictly followed (WHO, 2003). Similarly, in agar wel diffusion method, wells will be made on Muller Hinton agar plates using a sterile cork borer of appropriate diameter after which desired amount of the antimicrobial or the extract concentration to be tested will be dispensed into each well (Shahidi Bonjar, 2004). In dilution technique the antimicrobial substance diluted from its highest concentration to the lowest concentration of chemotherapeutic agent capable of preventing growth of the test organism, which gives the Minimum Inhibitory Concentration (MIC) (Abdelraouf, 2009).

# **1.5.** Plant materials as treatment for gastrointestinal diseases

Infections associated with bacterial pathogens are among some of the indications for treatment with traditional remedies that include plant products (Njoroge and Busman, 2007). Medicinal plants have been reported to cure urinary tract infections, gastrointestinal disorders, respiratory diseases and cutaneous infections. For example, the essential oil and eugenol purified from Ocimum gratissimum has been reported to treat pneumonia, diarrhea and conjunctivitis (Nakamura et al., 1999). Plants are powerful remedies in the treatment of abdominal infections especially dysentery, diarrhea and gastroenteritis (Mthabe et al., 2006; Lategan et al., 2009). Several studies have shown that diarrhea and gastroenteritis are a major cause of mortality in children under 5 years in developing countries (Ouattara et al., 2013). Many researchers in Ethiopia have identified medicianl plants as a remedey for the treatment of diarreheal diseases and gastrointestinal disorders (Table 2) (Mesfin et al., 2003; Gidey and Samuel, 2005; Ermias et al., 2008; Haile et al., 2008; Fisseha et al., 2009; Nasir et al., 2011; Rainer et al., 2011; Balcha Abera, 2014; Elizabeth et al., 2015). In Bale zones, several traditional medicinal plants have been also utilized to treat diarreheal disease and gastrointestinal disorders among which C. aurea, V. amygdalina, and R. nepalensis are the main and frequently usable (Table1and 2) (Ermias et al., 2008; Nasir et al., 2011; Rainer et al., 2011).

# 1.5.1. Calpurnia aurea

Studies have indicated that the plant *Calpurnia aurea* (wild), locally called Cheketa or cheekaa in Afan Oromo and Digita in Amharic language belongs to the family Fabaceae found almost in all areas of Bale zone especially in Mena Angetu forest (Ermias *et al.*, 2008; Nasir *et al.*, 2011). Decoction of leaf, seed and root of this plant are used as medicine in Bale zone as well as in different areas of the country to treat diarrhea, bloody diarrhea and abdominal pain (Gidey and Samuel, 2005; Ermias *et al.*, 2008; Nasir *et al.*, 2011; Abiyu *et al.*, 2014; Elizabeth *et al.*, 2014; Getaneh *et al.*, 2015).

Table 1: Common medicinal p	plants of Bale zone used to treat	gastrointestinal problems
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	Plant's Name		Plant	Disease	Study	
		Local Name	part	treated	area in	
R/		(Afan	used		Bale zone	
Ν	Scientific name	Oromo)				Reference
1	Hypericum revolutum	Garramba	L			
2	Cucumis ficifolius	Han'chote	R	Stomach		
3	Croton macrostachyus		В	problems		
				Intestinal		
4	Ocimum sp.	Damakase	L	infections	Odo -Bulu	
5	Allophylus abyssinicus	Habarra	F	Stomach	Demaro,	(Rainer et
6	Phytolacca Dodecandra	Endode	R	problems	Bale	al., 2011)
7	Asystasia guttata	Baquli	R	Diarrhoea		
8	Barleria argentea	Hokto	R&L	Diarrhoea		
9	Blyttia fruticulosum	Joshani	В	Diarrhoea	Mana	(Ermias et
10	Calpurina aurea	Chekata	L	Diarrhoea	Angetu	al., 2008)
						(Nasir et al.,
11	Calpurina aurea	Chekata	R&B	Dysentery	Bale zone	2011)
10		D 1. 1		D: 1	Mana	(Ermias <i>et</i>
12	Cissampelos pareira	Baltoke	R	Diarrhoea	Angetu	<i>al.</i> , 2008)
10			D	Vomiting,	Mana	(Ermias <i>et</i>
13	Cissus populnea	Gangalto	R	Diarrhea	Angetu	al., 2008)
14	Colocassia esculenta	Gondire	L	-	M	
15	Convolvulus siculus	LeIbab	L	-	Mana	(Ermias $et$
16	Dioscorea quartiniana	Gishu	R		Angetu	al., 2008)
17	Hibiscus luduwigii	Bulanbula	R	Vomiting		
18	Kleinia abyssinica	Burka	R	and		
19	Lippia javanica	Sukahi	L	Diarrhea	M	
20	Linnia ado maia	Sukahi	L	Abdominal Irritation	Mana Angetu	(Ermias <i>et al.</i> , 2008)
20	Lippia adoensis Oleae Cuspidata	Ejersa	L	IIIItatioli	Angetu	<i>u</i> ., 2008)
21	Olinia rochetiana	Guna	L	-		
23	Pentas lanceolata	Dhumuga	L	-		
23	Pittosporum viridiflorum	Ara	B	Stabing	Mana	(Ermias <i>et</i>
25	Pterolobium stellatum	Kejima	B	Pain	Angetu	(Erimas et al., 2008)
26	Rhynchosia densiflora	Rejillia	R	Vomit and	Mana	(Ermias <i>et</i>
27	Rumex nepalensis	Shabbe	R	Diarrhoea	Angetu	(Linnas <i>ei</i> <i>al.</i> , 2008)
27		Shucce	R	Diamoca	Odo Bulu-	<i>un</i> , 2000)
				Stomach	Demaro,	(Rainer et
28	Rumex nepalensis	Shabbe	R	Problem	Bale	al., 2008)
				Abdominal		(Meseret &
29	Rumex nervosus	Umbaco	L	pain	Gololcha	Egigu,2014)
					Mana	(Ermias <i>et</i>
30	Solanum giganteum		R	Diarrhoea	Angetu	al., 2008)
31	Stephania abyssinica	Kalala	R	_		7
32	Talinum caffrum	Burka	R	Stabing	Mana	(Ermias et
33	Thalictrum rhynchocarpum	Bala/Sirebizu	R	Pain	Angetu	al., 2008)
				Anti-		(Nasir et al.,
34	Vernonia amygdalina		L	dysentery	Bale zone	2011)
					Mana	(Ermias et
35	Zinnia peruviana		R	Diarrhea	Angetu	al., 2008)
				~ .		(Meseret &
26	411.			Stomach	G 1 1 7	Egigu,
36	Alliums sativa	Qullubi adii	Bu	ache	Gololcha	2014)
37	Rubia cordifolia	Anqis	R	Diarrhoea	Bale Mou.	Haile <i>et al.</i> ,
38	Tagetes minuta	Hada Gola	L	Diarrhoea	Nati. Park	2008)

	Plant's Name		Plant	Disease	Study	
R/ N	Scientific name	Local Name (Afan Oromo)	part used	treated	area in Bale zone	Reference
IN		Oromo)		Stabbing	Bale Mou.	Haile <i>et al.</i> ,
39	Gladiolus dalenii	Kelede	R	Pain	Nati. Park	2008)
40	Dovyalis abyssinica	Koshimo	W	Diarrhea	Bale Mou. Nati. Park	(Haile <i>et al.</i> , 2008)
41	Lippia adoensis	Sukayee	L	Diarrhea		
42	Solanum incanum	Hiddi	R	Stomach	Gololcha	(Meseret & Egigu,2014)
43	Solanum macracanthum	Hiddi	R	health and sudden pain	Odo Bulu and Demaro	(Rainer <i>et al.</i> , 2011)
				Abdominal	Bale Mou.	(Haile <i>et al.</i> ,
44	Solanum adoens	Hiddi	L	irritation	Nati. Park	2008)

Key: Plant parts: Bark (B), Bulb (Bu), Flower (F), Leaf (L), Root (R), Seed (S) and Whole: plant part (W)

# 1.5.2. Vernonia amygdalina

Vernonia amygdalina commonly called bitter leaf is a homely plant that grows almost everywhere especially in the tropical areas of Africa which belongs to the family Astaraceae. It is regarded as a wonderful gift from God to mankind because of its numerous medicinal values including cure for stomach ache, skin infections, diabetes, insomnia, tooth ache, acne, pneumonia, stoke, arthritis, fatigue, cough and bleeding (Agbogidi and Akpomorine, 2013). The plant locally named Girawa or Ebicha (Amharic and Afan Oromo language respectively) has a remarkable role as anti dysentery in Ethiopia particularly in Bale's society (see table 2) (Fisseha et al., 2009; Nasir et al., 2011; Abiyu et al., 2014; Gemedo et al., 2014; Genene and Reddy, 2015).

# 1.5.3. Rumex nepalensis

The genus Rumex comprises of about 200 species of herbs. R. nepalensis belongs to the family Polygonaceae is locally known as "Tult and Shabbe", Amharic and Oromic, respectively. It grows abundantly in many parts of the country. The use of *R. nepalensis* for various therapeutic purposes is well known in Ethiopian traditional medicine, especially in Bale zone. The pounded root is given to humans in case of diarrhea, stabbing pain, abdominal colic and stomach problems (Table 2) (Mesfin *et al.*, 2003; Ermias *et al.*, 2008; Rainer *et al.*, 2011; Abiyu *et al.*, 2014; Balcha Abera, 2014; Elizabeth *et al.*, 2014; Getnet *et al.*, 2015).

	Plant's N		Plant			
	Scientific		part			
Family	name	Local Name	used	Disease treated	Study area	Reference
		Chekata (Oro.)	L	Diarrhoea/ Bassa	Mana Angetu	(Ermias et al., 2008)
				Stomachache and		
		Chekata (Oro.)	R &B	dysentery	Bale zone	(Nasir et al., 2011)
		Digita (Amh.)	F	Abdominal pain	Fiche	(Abiyu et al., 2014)
				Child with		(Elizabeth et al.,
Fabaceae	a	Digita (Amh.) Zikita (Amh.)	L	diarrhoea (tekmat)	Fiche	2014)
ace	ure			Diarrhea &		
Fab	ai	Zikita (Amh.)	S	Bilharziasis		(Getnet et al., 2015)
	0	Zigita (Amh.)	R	Bloody diarrhea	Libo Kemkem	(Getnet et al., 2015)
				Vomiting and St-		(Gidey and Samuel,
			R& B	omach problem	Gindeberet	2005)
		Digita (Amh.)	L	Diarrhea	West Gojjam	(Getaneh et al., 2014)
			L	Anti-dysentry	Bale zone	(Nasir et al., 2011)
0	a	Girawa (Amh.)	L	Abdominal pain	Fiche	(Abiyu et al., 2014)
Asteraceae	din	Girawa (Amh.) Ebicha (Oro.)	L	Diarrhea	A/Negelle	(Gemedo et al., 2014)
rac	gdd					(Genene and Reddy,
ste	. · · · · · · · · · · · · · · · · · · ·	Ebicha (Oro.)	L	Diarrhea	Guji	2015)
A	V. an	Ebicha ()	L	Diarrhea	SNNPR	(Fisseha et al., 2009)
		Shabbee (Oro.)	R	Diarrhea	Mana Angetu	(Ermias et al., 2008)
					Odo Bulu-	
	is	Shabbee (Oro.)	R	Stomach problems	Demaro, Bale	(Rainer et al., 2011)
				Stomach		
	R. nepalensis	Tult (Oro.)	R	problems	Gimbii	(Balcha Abera, 2014)
5 D	pal			Stabbing pain,		
ace	ne	Lut (Amh.)	R	Diarrhea	Fiche	(Abiyu <i>et al.</i> , 2014)
Polygonacea				Tonsillitis and		
lyg		Tult (Amh.)	R	diarrhea	Libo Kemkem	(Getnet <i>et al.</i> , 2015)
Pol		Muca-araba				
		(Oro.)	R	Abdominal colic	Jimma	(Mesfin <i>et al.</i> , 2003)
				stomach ache		(Elizabeth <i>et al.</i> ,
IZ D1		Tult (Amh.)	R	(megagna)	Fiche	2014)

Table 2: Ethno-medicina	luce of the three sele	ated medicinal plant	in different	areas of the country
Table 2. Eulilo-medicina	i use of the three sele	cieu meulemai piana		areas of the country

**Key:** Plant parts: Bark (B), Flower (F), Leaf (L), Root (R) and Seed (S); Study area: Southern Nations, Nationalities and Peoples Regional State (SNNPR) Plant local names: Oromic language (Oro.), Amharic language (Amh.)

# 1.6. The Test Organisms

# 1.6.1. Escherichia coli

The genus of the bacterium *E. coli* is named for a scientist, German Pediatrician Theodor Escherich, whereas its specific epithet, coli, reminds us that *E. coli* live in the colon, or large intestine. He isolated the bacterium in 1885 from feces of infants with diarrhea. Later on, it was observed that the organism is a commensal of gastrointestinal tract and pathogenic and commensal strains can be distinguished by serotyping. Enterohaemorrhagic strains of *E. coli* have emerged as important enteric pathogens in recent years (Gerard *et al.*, 2010). *E. coli* is a gram-negative bacterial pathogen that causes many infectious diseases in human such as urinary tract infections and common intestinal diseases. Especially Enteropathogenic *Escherichia coli* (EPEC) that adheres to intestinal epithelial cells, causing diarrhea and constitutes a significant risk to human health that remains an important cause of infant mortality in developing countries (Matias *et al.* 2011).

Due to the ease of access of pathogens ingested with food, the human gastro-intestinal tract is susceptible to diarrhoeagenic *E. coli* infections. Several *E. coli* pathogens have been implicated with diarrhoeal illness, a major public health problem worldwide, with over 2 million deaths occurring each year (www.who.int); The major categories of diarrhoeagenic *E. coli* strains include enteropathogenic *E. coli* (EPEC), enterotoxigenic *E. coli* (ETEC), enteroinvasive *E. coli* (EIEC), enterohemorrhagic *E. coli* (EHEC) enteroaggregative *E. coli* (EAEC), and diffuse adhering *E. coli* (DAEC) (Huiwen *et al.*, 2005). As briefly described in table 3 each of these diarrhoeagenic *E. coli* strains characterized by their own diarrheal disease and how human beings get infected by

disease	they	develop.
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Table 3: Diarrhoeagenic groups of Escherichia

Group Designation	Disease, Site of Infection	Pathogenic Mechanism	
Enteropathogenic (EPEC)	Watery diarrhea, small intestine	Adherence	
Enteroinvasive (EIEC)	Watery diarrhea followed by bloody diarrhea, colon	Invasion	
Enterotoxigenic (ETEC)	xigenic (ETEC) Watery diarrhea, small intestine		
Shiga-toxin producing (STEC) Alsoknown as:EnterohemorrhagicVerotoxigenic (VTEC)	Watery diarrhea, sometimes bloody, especially for O157:H7, colon	Shiga toxins, type 1 and/or type 2	
Enteroaggregative (EaggEC)	Watery diarrhea, small intestine and colon	Adherence	
Diffuse adhering (DAEC)	Watery diarrhea, small intestine	Adherence	

Source: Emanuel and Lorrence (2009)

# 1.6.2. Staphylococcus aureus

Staphylococcus aureus (a gram-positive coccus) was discovered in 1880 by the surgeon Sir Alexander Ogston. He observed grape-like clusters of bacteria when examining a purulent discharge from patients with postoperative wounds during microscopy. He named them staphylé, the Greek expression for a bunch of grapes. In 1884, Rosenbach succeeded in isolating yellow bacterial colonies from abscesses and named them *S. aureus*, "*aureus*" from the Latin word for golden. Colonization with *S. aureus* is an important risk factor for subsequent *S. aureus* infection (Wertheim *et al.*, 2004; Lowy, 1998).

Staphylococcus aureus belongs to the family Micrococcaceae and is part of the genus Staphylococcus, which contains more than 30 species such as *S. epidermidis, S. saprophyticus* and *S. haemolyticus*. Among the staphylococcal species, *S. aureus* is by far the most virulent and pathogenic for humans. *Staphylococcus aureus* is a 1 $\mu$ m, Gram-positive cell that in the laboratory may be observed as single cells, in pairs or as grape-like irregular clusters. It is characterized as coagulase negative and catalase positive, non-motile, non-spore-forming and as facultative anaerobic (Washington, 2006).

*Staphylococcus aureus* is generally thought of as an extracellular pathogen, but it can be internalized by a variety of cell types *in vitro* (eg, fibroblasts, osteoblasts, keratinocytes, and endothelial cells (Hauck C, 2006). *Staphylococcus aureus* causes a wide range of infections from a variety of skin, wound and deep tissue infections to more life-threatening conditions such as pneumonia, endocarditis, septic arthritis and septicemia. In addition, *S. aureus* may also cause food poisoning, scalded-skin syndrome and toxic shock syndrome, through production of different toxins (Washington, 2006).

*Staphylococcus aureus* usually grows on the nasal membranes and skin; it also is found in the gastrointestinal and urinary tracts of warm-blooded animals which is a major cause of food poisoning caused by ingestion of improperly stored or cooked food in which *S. aureus* has grown (Prescott *et al.*, 2008).

*Staphylococcus aureus* is very resistant to heat, drying, and radiation; it is found in the nasal passages and it is also a frequent cause of skin lesions on the hands. From these sources it can readily enter food. If the bacteria are allowed to incubate in certain foods, they produce heat-stable enterotoxins that render the food dangerous even though it appears normal. Once the bacteria have produced the toxin, the food can be extensively and properly cooked, killing the bacteria without destroying the toxin. Intoxication can therefore result from food that has been thoroughly cooked. Thirteen different enterotoxins have been identified; enterotoxins A, B, C1, C2, D, and E are the most common. These toxins appear to act as neurotoxins that stimulate vomiting through the vagus nerve (Prescott *et al.*, 2008; Gerard *et al.*, 2010).

Typical symptoms include severe abdominal pain, cramps, diarrhea, vomiting, and nausea. The onset of symptoms is rapid (usually 1 to 8 hours) and of short duration (usually less than 24 hours). Prevention and control involve avoidance of food contamination, and control of personnel responsible for food preparation and distribution (Prescott *et al.*, 2008). Nearly any food can be contaminated with *S. aureus*, but those with a starch or cream base are most likely candidates. Cream pies, dairy products, poultry products, and picnic foods such as potato salad are common culprits. Contamination is difficult to detect because it produces no change in the food's appearance, taste, or odor (Jacquelyn, 2008).

## 1.6.3. Salmonella typhi

Salmonella is the causative agent of salmonellosis. It is a rod-shaped gram-negative facultative anaerobe bacterium belonging to the Enterobacteriaceae family. Among more than 2,300 closely-related Salmonella serovars recognized, S. typhi and Paratyphi are pathogenic exclusively for humans, and cause systemic infections and typhoid fever (McClelland et al., 2001). Salmonellosis is more prevalent in developing parts of the world in Africa, Asia, and South America.Typhoid fever is an acute, life-threatening febrile illness caused by the bacterium S. typhi and Paratyphi, and there are estimated 20 million cases and 200,000 deaths worldwide each

## year (Crump et al., 2004).

Salmonella organisms penetrate the mucosa of both small and large bowel, coming to lie intracellularly where they proliferate. After ingestion in food or water, typhoid organisms pass through the pylorus and reach the small intestine. They rapidly penetrate the mucosal epithelium via either microfold cells or enterocytes and arrive in the lamina propria, where they rapidly elicit an influx of macrophages (Mp) that ingest the bacilli but do not generally kill them. Acute typhoid fever is characterized by prolonged fever, disturbances of bowel function (constipation in adults, diarrhoea in children), headache, malaise and anorexia (WHO, 2003).

#### 1.6.4. Pseudomonas aerugenosa

Pseudomonas aeruginosa is a motile, non-fermenting bacterium, Gram-negative organism belonging to the family Pseudomonadaceae, opportunistic pathogen capable of infecting humans with compromised natural defenses and causing severe pulmonary disease. It is one of the leading pathogen associated with nosocomial (hospital-acquired, infections are those not present or incubating at the time of hospital admission, but usually develop post-admission infections) (Alaa Alhazmi, 2015). Pseudomonas aeruginosa is an opportunistic pathogen familiar to every hospitalist and intensivist. It is a frequent cause of nosocomial infections, many of which are responsible for substantial mortality even when treated with appropriate antibiotics (Juhas et al., 2005).

## 1.6.5. Shigella dysentery

Shigella organisms are a group of gram negative, facultative intracellular pathogens. They were recognized as the etiologic agents of bacillary dysentery or shigellosis in the 1890s. Shigella were discovered over 100 years ago by a Japanese microbiologist named Shiga, for whom the genus is named. Shigella was adopted as a genus in the 1950s. These organisms are members of the family Enterobacteriaceae and tribe Escherichieae; they are grouped into 4 species: Shigella dysenteriae, Shigella flexneri, Shigella boydii, and Shigella sonnei, also known as groups A, B, C, and D, respectively (Gomez et al., 1997).

## 1.7. Extraction Solvents

Several investigators on medicinal plants have indicated that organic solvents such as alcohols are extensively used for crude extraction before being re extracted to obtained purified active compounds using some other organic solvents (Francis, and Richard, 2007; Anibijuwon *et al.*, 2012). Polar and ionic solutes are easily dissolved in the polar solvents. When both of the solutes and the solvents are polar, they have attraction between each other (Francis, and Richard, 2007). Solvents are generally classified by their polarity, and considered either polar or non polar, as indicated by the dielectric constant (Table 4). The greater the dielectric constant, the greater the polarity. Solvents with dielectric constants greater than about 5 are considered "polar" and those with dielectric constants less than 5 are considered "nonpolar" (*http://chemwiki.ucdavis*). Plant extracts are expected to be polar compounds which indicate that they dissolve in polar solvents and results clearly showed that utilization of polar solvent enable extraction of significant amounts of plant extracts (Farooq and Roman, 2012). Polar protic solvents, especially water (78.5) and methanol (32.6) have larger value of polarity, hence they can dissolve plant materials better than the other solvents (Francis, and Richard, 2007).

Generally higher extract yields, phenolic contents and plant material antioxidant activity were obtained using aqueous organic solvents, as compared to the respective absolute organic solvents (absolute methanol and aqueous methanol, i.e., methanol: water, 80:20) (Bushra *et al.*, 2009). Methanol shows both polar and non-polar characteristics. Consequently, these characteristics give methanol the unique ability to be used in an extensive range of applications (Esteban *et al.*, 2006).

Table 4: Common solvents used in organic chemistry

		Dielectric				
Solvent	<b>Boiling Point, Celsius</b>	Constant				
Non-Polar Solven	Non-Polar Solvents					
Pentane, $C_5H_{12}$	36	1.8				
Hexane, $C_6H_{14}$	69	1.9				
Benzene, C <sub>6</sub> H <sub>6</sub>	80	2.3				
Chloroform, CHC <sub>13</sub>	61	4.8				
Diethyl ether, (CH <sub>3</sub> CH <sub>2</sub> ) <sub>2</sub> O	35	4.3				
1,40-Dioxane,Cyc- (CH <sub>2</sub> CH <sub>2</sub> OCH <sub>2</sub> CH <sub>2</sub> O)	101	2.3				
Polar Parotic Solve	nts					
Water, H <sub>2</sub> O	100	78.5				
Methanol, CH <sub>3</sub> OH	65	32.6				
Ethanol, CH <sub>3</sub> CH <sub>2</sub> OH	78.5	24.3				
isopropyl alcohol, CH <sub>3</sub> CH(OH)CH <sub>3</sub>	82	18				
acetic acid, CH <sub>3</sub> COOH	118	6				
Polar Aprotic Solve	ents					
Dichloromethane, CH <sub>2</sub> Cl <sub>2</sub>	40	9.1				
tetrahydrofuran (THF), cyc - (CH <sub>2</sub> ) <sub>4</sub> O	66	7.5				
ethyl acetate, CH <sub>3</sub> C(O)OCH <sub>2</sub> CH <sub>3</sub>	77	6				
Acetonitrile, CH <sub>3</sub> CN	81.6	37.5				
dimethylformamide (DMF), HCON(CH <sub>3</sub> ) <sub>2</sub>	153	38				
dimethyl sulfoxide (DMSO), CH <sub>3</sub> SOCH <sub>3</sub>	189	47				
Acetone, CH <sub>3</sub> COCH <sub>3</sub>	56.5	21				
hexamethylphosphoric triamde (HMPT), [(CH <sub>3</sub> ) <sub>2</sub> N] <sub>3</sub> PO	232	30				

Source: (Francis and Richard, 2007).

## CONCLUSIONS

This review stated that plants are valuable sources for new compounds and should receive special attention in research strategies to develop new antimicrobials urgently required in the near future. The importance of medicinal plants and traditional health systems in solving the health care problems of the world is gaining increasing attention. Because of this resurgence of interest, the research on plants of medicinal importance is growing phenomenally at the international level, often to the detriment of natural habitats and protects populations in the countries of origin. In Ethiopia since many years the peoples are using plants as the medicine. The plant contains various phytochemical, which would act on the pathogenic microorganism and hinder their growth. The scanty of researches are available on antibacterial activity of medicinal plants in Ethiopia, but still we need to explore many antimicrobial compounds from plants. These kinds of the research are baseline information for the development of new drugs, which is bench mark in science. Commercially many synthetic drugs are available in market, but microorganisms are resistant to many antibiotics. So research should need to develop new synthetic compound from the plant source.

## **Competing Interests**

The authors declare that they have no competing interests.

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