Antibacterial potential of *Tridex procumbens* L.against Human pathogen.

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I. INTRODUCTION

Plants and its parts are rich source of bioactive molecule which offers a new source of antibacterial agent. In recent years *Tridex procumbens* plants have been reported Antibacterial activity of the methanol, chloroform, ethanol, & aqueou extract (Suryakantet.al.2017). *Tridexprocumbens* is medicinal herb and used by many ethano-medical practitioners, In India *Tridex procumbens* is employed as indigenous medicine. It has been found to possess significant medicinal properties against Dysentery ,Diarrhoea, Stomach ache and Liver disorder. It also shows Antidiabetic and anti-inflammatory actions. (Jain 2012). *Tridex procumbens* plant is found all over India .belonging to family-Asteraceae. It is commonly known**coat button**. It has been extensively used in Indian traditional medicinal system. The all parts of this plant contains number of secondary metabolite and have positive effect on pathogenic bacteria. (VeenaGayathri *etal*2015). *Tridex procumbens* extract have been reported antibacterial activity against gram positive organism. (V.Bharathi *etal* 2012) The present studyhas been undertaken to establish the antibacterial potential of *Tridex procumbens*. However, thewide-spread use of this methodology for antimicrobial susceptibility testing required flow spectrophotometer equipment in various laboratories.

II. MATERIAL AND METHOD

- 1. **Collection of plant material**: The fresh plant of *Tridex procumbens* were collected from different sites of Aurangabad and local village.
- 2. **Methanol extract:** 40 gm powder of fresh and .shad dry Root stem and leaves extracted by Soxhlet extraction process.
- 3. **Test organism**: The authentic culture of human pathogenic bacteria viz. Salmonella typhimurium ,*Pseudomonas aeruginosa, Shigella flexneri, E. coli and Staphylococcus aureus* were obtained from the department of Microbiology, Deogiri College,Aurangabad, Maharashtra. In vitro antibacterial assay of plant and fungal extract was carried out by using 96- well plate method.
- 4. 96 well plates method: About 100μl sterile Mueller-Hinton broths medium was loaded into each well along with 2μl serial diluted human pathogenic bacteria suspension, next 2,4, 6, 8, and 10μl concentrations of methanol extracted plant and ungal extract was added to each well of 96- well plate. Control was prepared by nutrient broth and bacterial suspension without adding extract. The prepared experimental 96- well plate was sealed with parafilm and incubated in incubator at 37°C for 24 hours. Finally optical density (OD) at 540nm was measured on the soectrophotometer of each sample (Ataee, *et al.*, 2012)

III. Results

Minimum inhibitory concentration (MIC) of the methanol extract was evaluated by 96 well plate method followed by optical density at 450 nm was measured among the 5 serial dilution for each pathogenic bacteria were tested. In which root, leaves and stem showed significant activity.2 µl concentration of root extract for *Salmonella typhimurium*, *Pseudomonas aeruginosa* and *Shigellaflexneri* was most significant.4µl concentration was effective for P. *aeruginosa* evaluating antibacterial activity also showed MIC at .6µlconcentration and .10µlconcentration.(Table –I).The stem of *Tridax procumbens* were showed effective control at 2,4,6,8 and 10 µlrespectively for *E.coli*, *Shigellaflexneri*, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Salmonella typhimurium*.(*Table –II*).*MIC for leaf extract was* 2,4,10 µl for ,*Shigella Flexneri*,

Pseudomonas aeruginosa,E.coli and *Salmonella typimurium.(Table- III)* Increase Concentration of root extract inhibits growth of bacteria. S.arueus shows more inhibition as compare to other bacteria.Increase and decrease Concentration of root extract inhibits growth of *S. typhi* bacteria.(table- I) similar results was observed in *E.coli*.Increase Concentration of stem extract was most effective for *Salmonella typimurium Shigella flexneri.(table-II)*. Leaf extract10µl *Salmonella typimurium* 2 µl for *Shigella flexneri. (table III)*

IV. DISCUSSION

Plants are a promising tool for pharmaceutical science. The review describes information of production of useful secondary metabolite and antibacterialactivity. (VeenaGyatri, Krishaswamy et al 2015) In this study Methanol extracted of plant was prepared. All extract of Root, Stem and,Leaves wereused for antibacterial test by 96 well plate methods. (Ataee, et al 2012)Control was prepared by nutrient broth and bacterial suspension without adding extract .According to optical density at 540 nm it was observed that Root extract 2 µl and 10 µl was most effective for S.typhi, S.flexneri, P. aurignos . 6 µl was most effective for E.coli.(Table- I) Stem extract of 10 µl was effective for S. typhi(Table- II) (Suryakant, Dr. SudhanshuDharDwivedi, SheerazAhmad 2017) methanol leaf extract of Tridex procumbenshows best result against Pseudomonas, and Klebsiella sp. According to(R. Dhanabalan, A. Doss, etal 2008) methanol extracts of this plant was investigated by agar disc and well-diffusion method against bovine mastitis causing Staphylococcus aureusstrains. In present study leaf extract is not shown significant activity (.Ankitajainetal 2015)methanol and aqueous extract of the plant was effective against E. coli and Bacillus subtilis.(Ahmad Mir S'Mahmood Dar 2016) The ethanolic extract of this plant showed highest zone of inhibition against *Escherichia coli* and *Staphylococcus aureus*. The plant part extract of this plant showed highest zone of inhibition against Klebsielle pneumonia. A total of 5 pathogenic bacteria were incubated at 37°C for 24 hours and anti bacterial activity was determine .antimicrobial results that inhibits the pathogenic bacteria 2µl concentration of Methanol extract of root was most effective for Salmonella typhimurium, Pseudomonas aeruginosa and Shigella flexneri. 2µl concentration of stem were significant for *E.coli*.(Graph –A) This paper describe although the high potential of root and stem.

Sr.no	Tridex	Bacterial concentration(2µl)					
	root	S.typhi	S.flexineri	P.aurignosa	S.arueus	E.coli	
	extract						
1	2µ1	0.00	0.00	0.00	0.02	0.01	
2	4µ1	0.01	0.01	0.00	0.01	0.01	
3	6µl	0.02	0.00	0.02	0.01	0.00	
4	8µ1	0.01	0.02	0.01	0.05	0.01	
5	10µ1	0.00	0.00	0.01	0.00	0.00	
	MIC	2,10µl	2,6,10µl	2,4µl	10µ1	6,10µ1	

Table I Antibacterial activity of Tridex procumbens root extract

 Table II Antibacterial activity of Tridex procumbens stem extract

Sr.no	Tridex	Bacterial concentration(2µl)					
	stem	S.typhi	S.flexineri	P.aurignosa	S.arueus	E.coli	
	extract						
1	2µ1	0.07	0.07	0.07	0.09	0.06	
2	4µ1	0.08	0.09	0.07	0.08	0.09	
3	бµl	0.07	0.06	0.07	0.08	0.09	
4	8µ1	0.07	0.07	0.06	0.07	0.08	
5	10µl	0.04	0.06	0.08	0.09	0.11	
	MIC	10 µl	6,10µl	8 µl	8 µl	2 µl	

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Sr.no	Tridex	Bacterial concentration(2µl)						
	leaf	S.typhi	S.flexineri	P.aurignosa	S.arueus	E.coli		
	extract		-	_				
1	2µl	0.14	0.09	0.13	0.16	0.16		
2	4µl	0.15	0.12	0.16	0.17	0.14		
3	6µl	0.14	0.13	0.14	0.18	0.17		
4	8µl	0.20	0.20	0.20	0.19	0.20		
5	10µ1	0.13	0.13	0.18	0.14	0.16		
	MIC	10 µl	2 µl	2 µl	10 µl	4 µl		

Table III Antibacterial activity of Tridex procumbens leaf extract



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