# Valorization and Management of Biogas Slurry

By Satyawati Sharma

At

Int. Workshop on "Promotion of Biogas Upgrading and Bottling in India and European Union" on August 22 - 24, 2013 , IIT Delhi

## Introduction

• No. of Biogas plants (BGFP) installed in India= 4.31

million (household and community type)(MNRE,2013)

- Of these 95.81% functional.
- Huge quantity of BGS is being produced
- BGS = A byproduct produced during biogas production
- BGS is good O.M (free from weed seeds and pathogens, nutrient rich)

• But due to high water content ;

92-94% in cattle dung based BGS97-98% in food waste BGS, not being utilized optimally.

- Transportation/ storage biggest problem.
- Sun drying –common method ,nutrients (N,S etc.) lost.

Other applications of BGS: As biopesticide (Aphid, Beetel) pisciculture, animal feed (pig), dye absorption, mushroom cultivation (only *Pleurotus*) (Some references)
 ✓ Not much field work has not been done

## **Objectives (MNRE sponsored project)**

- 1. Technological interventions for enrichment of BGS during dehydration and packaging operations.
- 2. Development of protocols for utilization of BGS in liquid (algal biomass production), semi solid (vermi compost), and solid form (mushroom & organic manure).
- 3. Integration of protocols for developing the entrepreneurship package (Integrated Approach).

# **1.Characterization of BGS**

#### **Characteristics of Fresh, Sundried and One month stored slurry**

Parameters tested	BGS from Pur	BGS from Purana Farm House, Vasant kunj, N.Delhi					
	Fresh	Sundried * Stored for one month					
Cellulose (%)	29.44	28.75	20.75				
			1601				

✓ ~18.18% N lost during sun drying ,Also time consuming process

### ✓ ~9% N loss in storage for one month

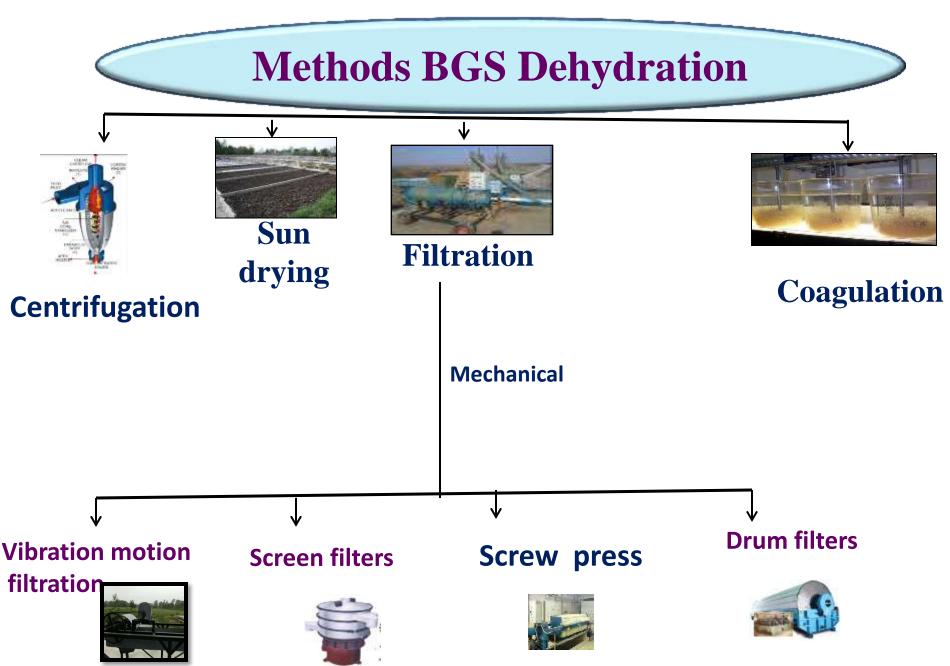
#### ✓ Also lot space in both the cases needed

C % (dry wt. basis)	37.25	38.5	16.99

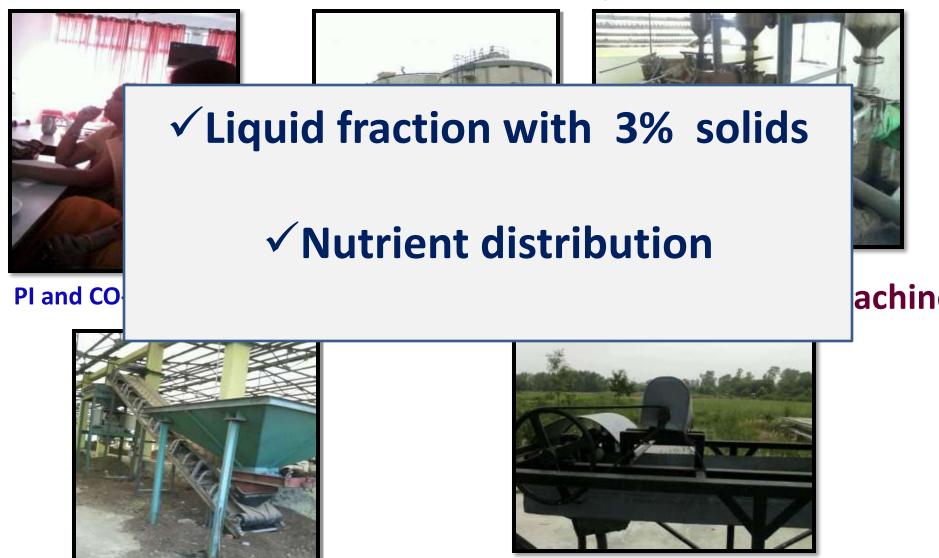
•1 inch thick layer of Slurry dried for 1 month

Nitrogen (TKN): 0.88%, Nitrate: 0.08%, Ammonical N: 0.07%

# **2. Dehydration of BGS**

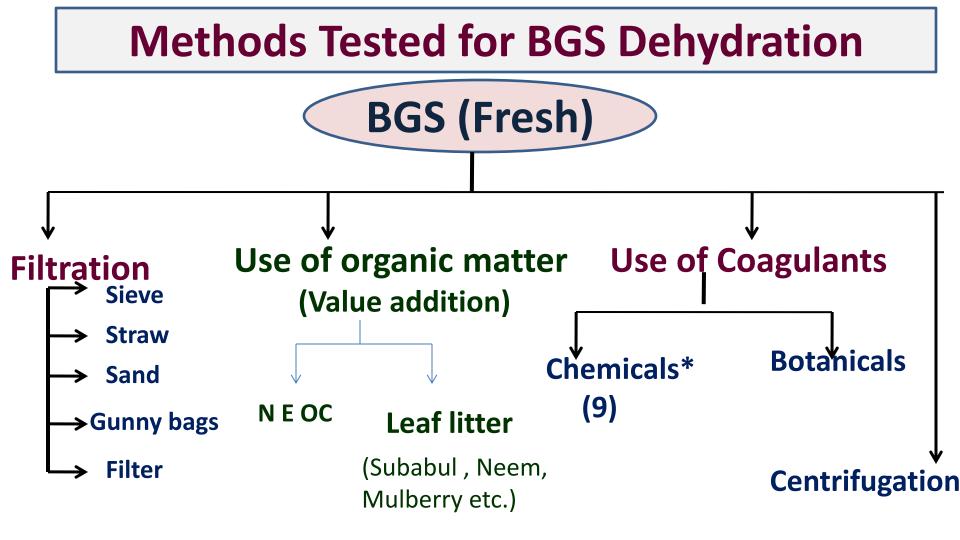


### Glimpses of Visit to PEDA installed / DSM operated Biogas Plant, Ludhiana, Punjab



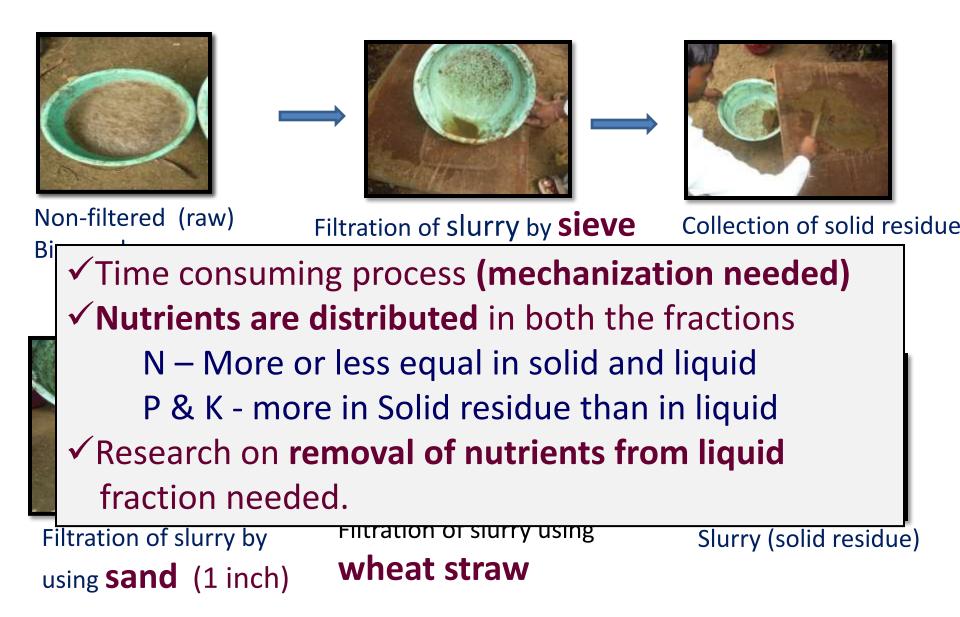
Solid and dried slurry separator

#### Vibration filtration unit at PAU, Ludhiana



\* Alum (with and without KOH,KCL), Iontosorb oxin, Polyhydroxamic acid (PHA), Polyacrylamide, Acrylamide, Gelatine, Epichlorohydrin, Chetosan, DMA

## **Filtration of Slurry**



#### **Filtration by Gunny Bags**

#### Fresh/Raw BGS



**At Micron** 



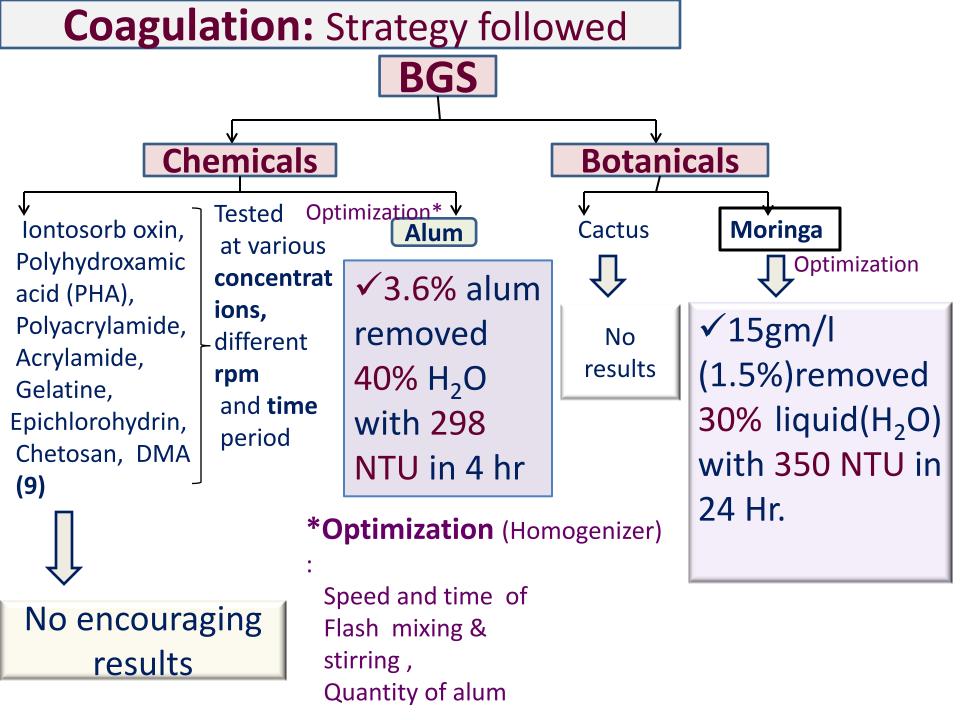






Water Obtained **Gunny Bag Filtration Set-Up** (~50 %) **Properties of removed water** EC: 5.83mS/cm , pH: 8.6, NTU: 812

Further work needed





Using Chemicals (KCl, KOH),

#### Polyelectrolytes

(Polyacrylamide, Chitosan,

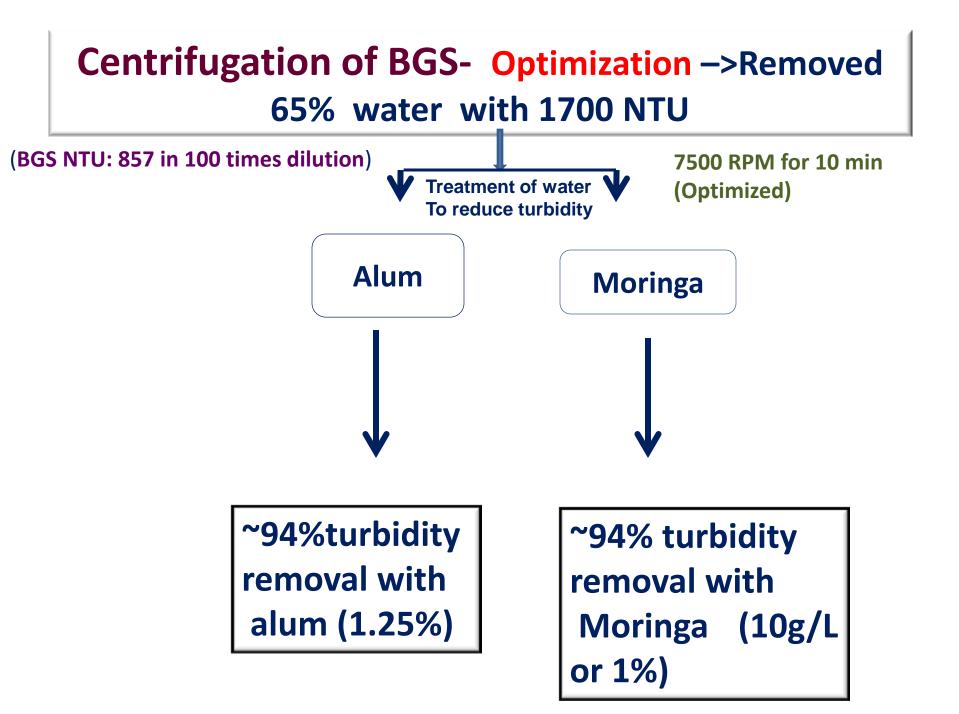
Epichlorohydrin, DMA)

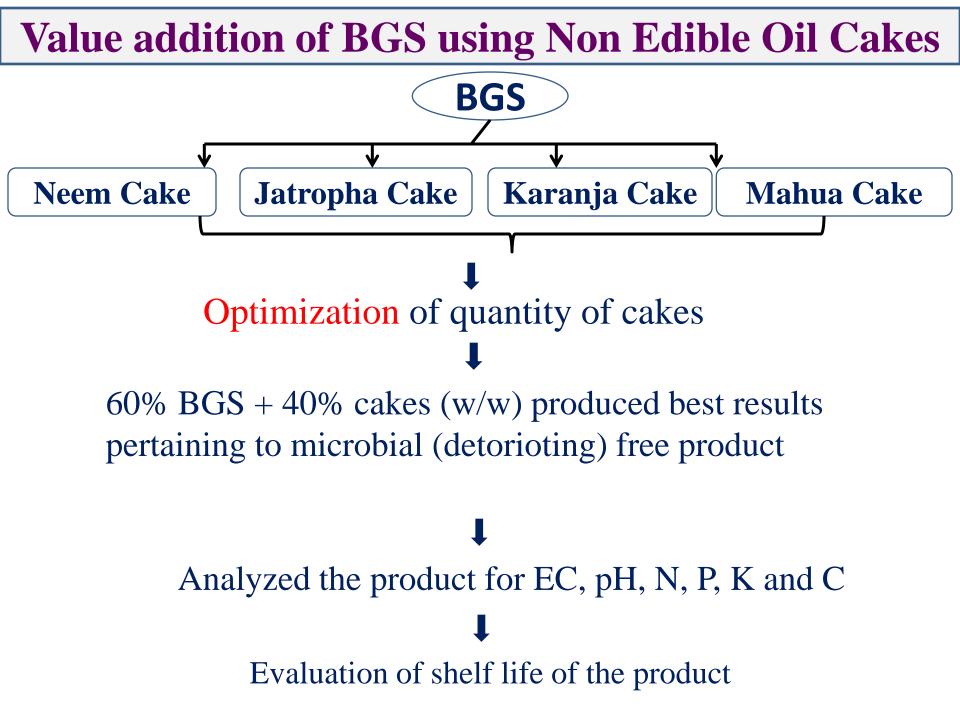
Best results with 2% alum and Dimethylamine (44% water removal with EC: 13.9 mS/cm; pH: 4.97 and NTU 242)

#### **Characteristics of water and Sludge separated by Moringa** (15g/L)

Parameter	Water	Sludge
EC (mS/cm)	1.4	8.3
рН	7.5	7.05
TOC (%)	15.8	64.5
N (%)	0.9	2.82
P%	0.5	1.51
K%	0.57	1.01

Quantity of water removed = 30% with 350NTU
 Enriched sludge/solid fraction(Protein in moringa seeds= 61%)





## **Properties of Value Added BGS (With NEOC)**

Sample	EC (mS/cm)	рН	TOC%	TKN%	C:N	%K	%P
Biogas Slurry	4.98	7.9	37.69	0.75	50.25	0.8	0.43
Mahua Cake	2.58	5.8	44.20	2.04	21.66	1.10	0.60
Mahua cake + BGS	3.71	6.39	40.78	1.86	21.92	1.08	0.56
(6:4)							
Karanja Cake	<ul> <li>✓ NPK content increased significantly</li> <li>✓ Work on optimum doses for different crops (seed</li> </ul>						0.90
Karana cake + l							1.38
(6:4)							
Neem Cake							1.1
Neem cake + B			•	•			1.38
(6:4)	germination)required						
Jatropa	2.41	6.01	44.35	5.03	7.35	1.2	1.41
Jatropha cake +BGS (6:4)	3.67	6.82	41.15	3.56	9.02	2.43	1.61

### Value Addition of BGS through Algae cultivation





Dried product

N(%) = 1.88

P(%) = 1.23

0.99

C(%) = 30.21

K (%)





Dried value added product

✓ Increased significantly from
initial value
(N= 0.88%; P= 0.58%; K= 0.87%)



BGS (1 L)+ Organic amendments +



#### Inoculated Chlorella (2% w/v) (0.9 x 10<sup>5</sup> cells / mL) After 25 days



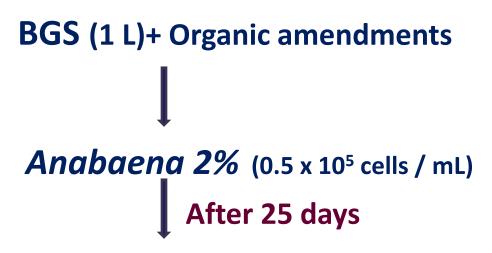
Growth of *C. minuttisima* in BGS with diff Conc. of Org Amend.

inuttisima

(40x)

Parameter	Control (Media)	BGS + Organi	BGS	0.0
		amend		
Chlorophyll a	1.21	1.60	-	Alter all shares and shares
Chlorophyll b	1.93	2.06	-	0
Total	2.91	3.66	-	0 0
Chlorophyll	N (%) =	1.35	✓ Increased	significantly from
Cell count (10 <sup>5</sup> cells/mL	P (%) =	1.11	initial value	
	K (%) =	1.12	(N= 0.88%; F	P= 0.58%; K= 0.87%)

## (C ) Anabaena





# Growth of *Anabaena* in BG S with diff conc. of amendment

Parameter	Control	BGS	+ BGS	6. 1 . C.	-
	(Media)	Orga	anic	A Com	and a
		ame	ndment		
Chlorophylla	<b>a</b> 0.41	0.55	-	5000 200	621:
Chlorophyll	h 1 3 1	1 46		South France & R	
Total	N (%) =	1.97	✓ Increased signi	ficantly from	
Chlorophyll	P (%) =	1.15	initial value		w of
Cell count	K (%) =	1.22	(N= 0.88%; P= 0.	50%; K = 0.07%	GS (best
(10 <sup>5</sup> cells/mL	.)			шеши, чолј	

# (D) Nostoc BGS (1 L)+ Organic amendments Nostoc (0.5 x 10<sup>5</sup> cells / mL) After 25 days

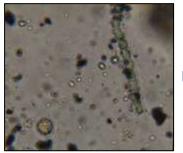


# Growth of *Nostoc* in BG S with diff conc. of amendment

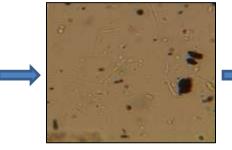
Parameter	Contro	BGS	+ BGS	
	(Media	) Orga	nic	40 00 00 00 00 00 00 00 00 00 00 00 00 0
		ame	ndment	· And Constants
Chlorophyll *	<b>'a</b> 0.38	0.43	-	ere and
Chlorophyll I		1 20	_	Code I Aller R. Goe
Total	N (%) =		✓ Increased sign	ificantly from initial
Chlorophyll	P (%) =	1.12	value	
Cell count	K (%) =	1.20	(N= 0.88%; P= 0	.58%; K= 0.87%)
(10 <sup>5</sup> cells/mL)	)			(*Micro gm/ml of medium)

### **BGA cultivated in BGS**

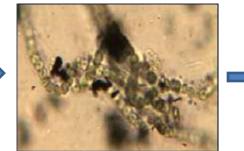
(All under 40X Phase Contrast Microscope)



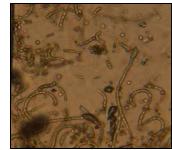
Anabaena (0 day)



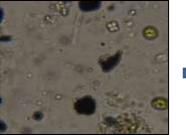
Anabaena (5th day) Budding cells



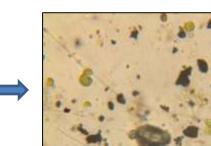
Anabaena (10th day)



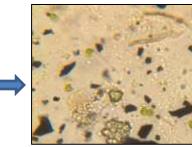
Anabaena (20th day) At 10X



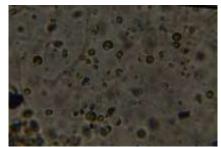
Chlorella (0 day)



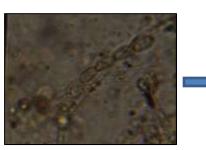
*Chlorella* (5th day) Cell division



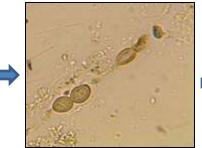
Chlorella (10th day)



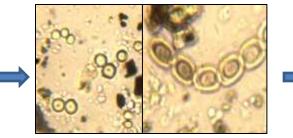
Chlorella (20th day)



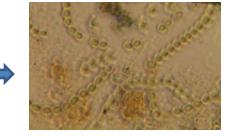
Nostoc



*Nostoc* (5th day) Cell division



Nostoc (10th day) Developing filaments



Nostoc (20th day)

# **BGS as CO-Composting Material**

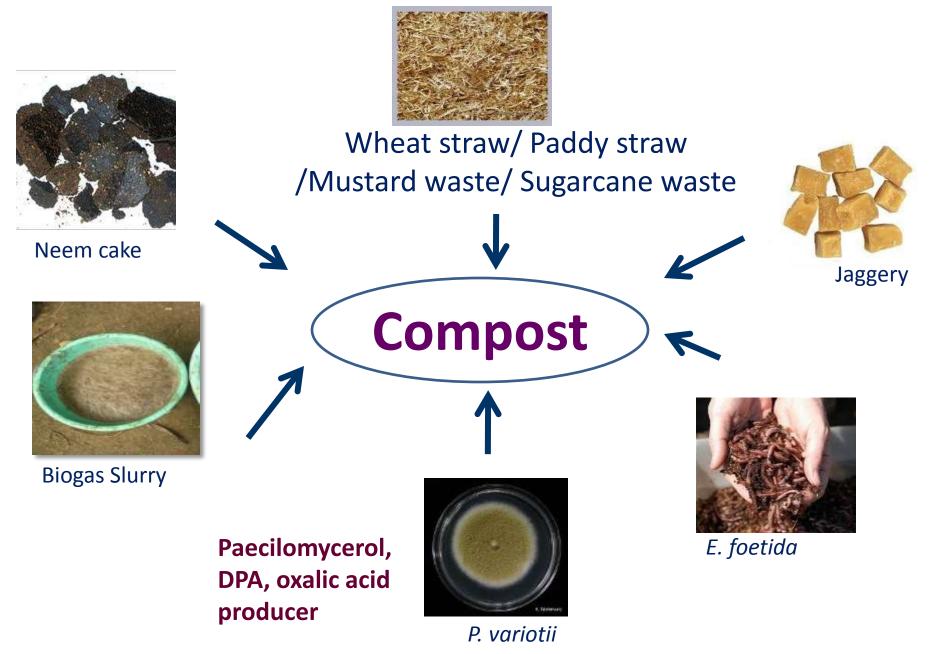


### Composting with Biogas Slurry in Micromodel, IITD





# **Materials used in Composting**



#### Use of Biogas Slurry in Composting using Wheat Straw

Substrate
 Combination
 BGS :Substrate

4:1 (Wheat straw)

2:1 (Paddy straw)

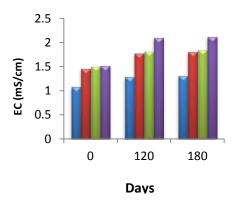
2:1( Mustard waste)

3:1 (Sugarcane baggase)

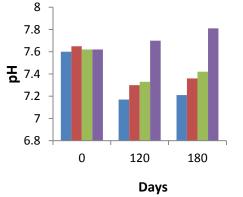
for ~ 65% moisture

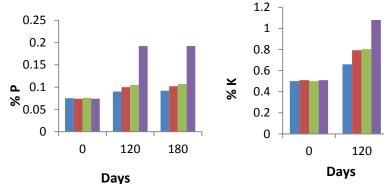
Treatments	Substrate combination for composting
T1	Straw + Slurry
T2	Straw + Slurry + Culture
Т3	Straw + Slurry + Culture + Jaggery(0.5%) + Neem cake (0.1%)
T4	Straw + Slurry + Culture + Jaggery + Neem cake + Earthworm

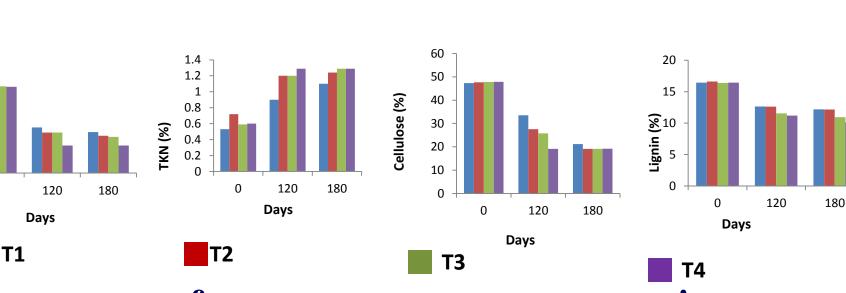
# Changes in parameters during composting of Waste (wheat straw) with BGS



TOC (%)

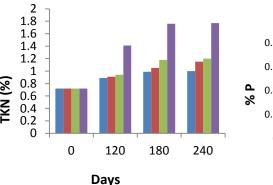


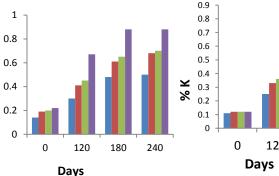


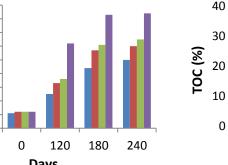


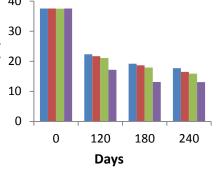
✓ Saturation in **four** months in T4 and more than **six** months in T1

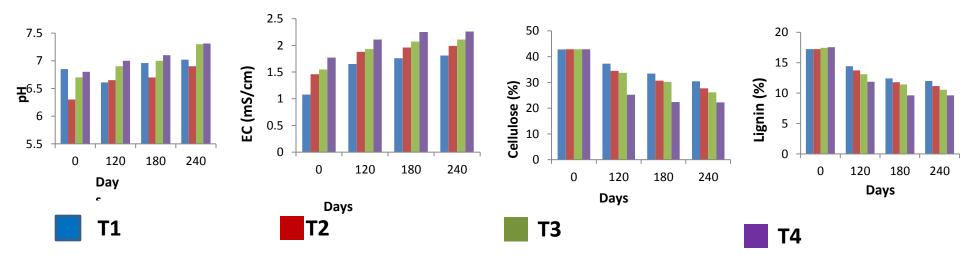
# Changes in parameters during composting of Waste (Paddy straw) with BGS











✓ Stabilization in T4 within Six months, in T1 in more than 8 months
 ✓ Similarly, Mustard waste: T4 = 5.5 months, T1= 7 months
 ✓ Sugarcane baggase: T4= 6 month, T1 >8 months

# **Use of BGS in Mushroom Cultivation**

• Types of Mushrooms Cultivated:

## 1.Pleurotus sajor-caju (Dhingri)

2. Agaricus bisporus (Button)

#### (200kg wheat straw+800 Lts BGS)











Button Mushroom Cultivation Using BGS in Micromodel,IITD

Ready for Distribution





Fructification

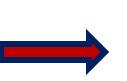


**Mushroom Harvest** 

**Fructification** 

#### Mushroom Cultivation at Shri Krishna Gaushala, Ghaziabad







#### Shri Krishna Gaushala, Ghaziabad

#### **Compost Production**



#### Starting of fructification (Shri Awasthi Ji, Gaushala)



#### Spawn Run

#### Mushroom production using BGS at Micromodel IITD









#### **Button Mushroom**

#### **Dhingri Mushroom**

### Yield of A. bisporus (Button) mushroom on BGS

- Experiments Conducted during November Feb.,2012 &2013 (Compost production=30days, Spawning +growth of mushroom fungi=30days)
- ≻Quantity of waste (wheat straw) used: 200 Kg &300kg
- ≻Quantity of compost formed: 125 kg &187.5kg.
- >Yield of mushroom: 24.09 kg/ 100 kg of compost in 2012
- >Yield of *Agaricus* mushroom : 20-22 Kg/100 kg of
- traditional compost

#### **Dhingri Mushroom Yield from Different Treatments of BGS**

BGS - straw combination	Mushroom yield (gm/kg dry substrate)	Biological efficiency (%)
100%	729	72.9
straw		
90%straw+10%	950	95
slurry		
80%straw+20%	700	70
slurry		
70%straw+30% slurry	600	60
60%straw+40% slurry	530	53
50%straw+50% slurry	230	23
25%straw+75% slurry	NIL	-

#### Yield of *Pleurotus florida* (Dhingri) on BGS with Karanja cake

S.No	Treatment	Yield (g/Kg)
1	Control	740.4
2	90% WS + 10% (Cake + BGS)	975
3	80% WS +20% (Cake + BGS)	760
4	70% WS +30% (Cake + BGS)	357
5	60% WS +40% (Cake + BGS)	Nil
6	50% WS +50% (Cake + BGS)	Nil

# Yield of *Pleurotus florida* (Dhingri) on BGS (from cattle dung mixed with mahua cake)

S. No	Treatment	Yield (g/Kg)
1	Control	700.36
2	90% WS +10% (Cake + BGS)	850.35
3	80% WS + 20% (Cake + BGS)	957.25
4	70% WS +30% (Cake + BGS)	686
5	60% WS +40% (Cake + BGS)	555.2
6	500% WS + 50% (Cake +	459.8
	BGS)	

#### Cake and BGS = 60:40

# Yield and Nutritional analysis of Mushroom fruit bodies cultivated on mahua cake based BGS

BGS mixed with straw combination	Proteins (%)	Total soluble sugars (%)	Fat (%)	Energy Kcal	P (mg/g)	K (mg/g )	Fe (ppm)
T1 (100% Straw )	29.6	32.33	2.01	1127.48	11.23	23.26	105.7
T2 (10%)	32.6	30.93	1.86	1149.2	11.56	23.46	129.6
T2 (20%)	33.42	31.06	1.826	1160.1	11.76	23.63	132.8
T2 (30%)	30.5	31.43	1.87	1122.3	11.35	23.06	134
T3 (10%)	36.23	29.06	1.84	1185	13.46	28.7	197.3
T3 (20%)	38.76	29.26	1.813	1220.26	15.0	29.86	200.6
T3 (30%)	34.61	29.46	1.826	1153.6	11.96	25.9	206.6

**T**<sub>1</sub> = 100% WS

 $T_2 = CD$  slurry control

T<sub>3</sub>= BGS plus DMC in 60:40

# Conclusions

# Filtration

Filtration distributed nutrients in both liquid as well as solid parts.

- ≻100% removal of solid (Colloidal ) particles not possible by sand, sieve, straw ,Screw press filtration.
- ➤Among all methods tried , gunny bag method was found suitable (>50% water removal) and can be employed at small biogas plants. However further work is needed.
- > Development of filtration unit using diff. mesh size sieves and motor (under progress)

# Centrifugation

- Centrifugation yielded 65% of water with turbidity 1700
  NTU.
- ➢Moringa and alum reduced turbidity from 1700 NTU to 99 and 97 NTU respectively.
- Moringa proved better than Alum..
   Sludge with 60% moisture vermicomposted with earthworms for 20 days yielded quality product rich in NPK.
- ➤Cost ?? involved in centrifugation (Total volume 500 L; Cake volume 20kg.; 3000 rpm) may be the limiting factor (~ 2.0 Lakhs).

# **Use of Coagulants:**

- Among all coagulants tested, alum (2%) along with Dimethylamine (DMA 0.016%) produced best results (44% water removal with EC 13.9 mS/cm and pH 4.97 in <4 hr)</p>
- The increase in pH (7-7.5) was possible with the use of KOH and lime but EC could not be decreased.
- ➢ Moringa seed powder removed 30% water, although it takes time (>12 hr), may be useful as it enhances the manurial value of sludge.

# Value Addition of BGS through BGA Cultivation:

Chlorella, Anabaena and Nostoc : The use of amendments in BGS improved the growth of BGA. The liquid BGS with growth of Chlorella, Anabaena and Nostoc can be used as liquid BGS based biofertilizers.

 Azolla along with BGS after 20 days growth produced the product with enhanced nutritional content (NPK). Quality Solid Biofertilizer.

➤Use of NEOC and Leafy litter: Not only reduced water content ,also improved quality

# **Mushroom cultivation:**

# **Pleurotus** :

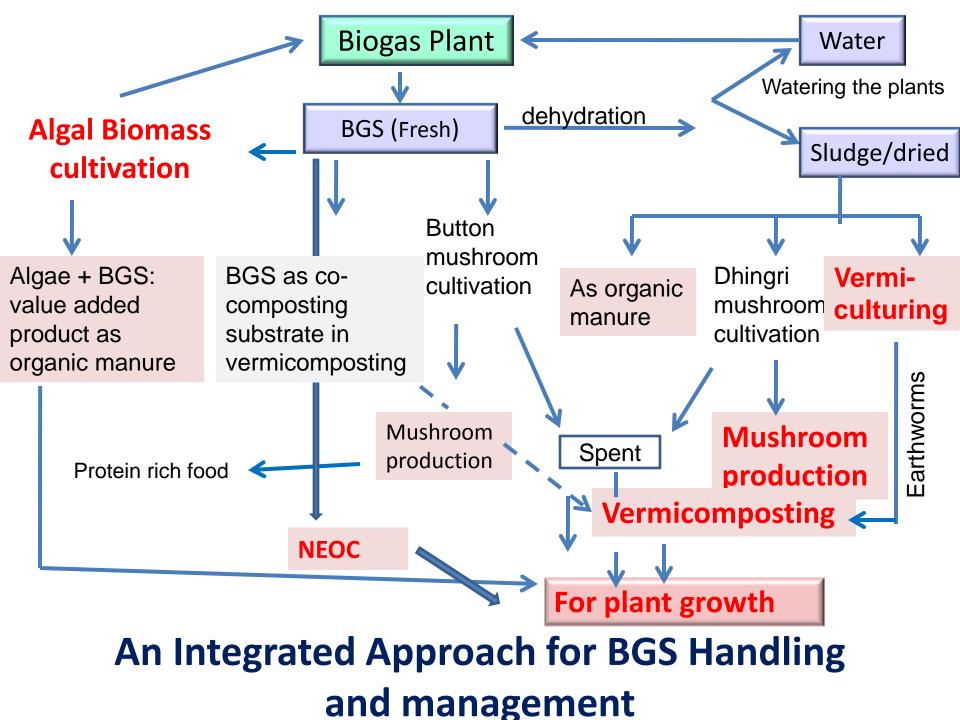
- ≻10% of BGS (amendment) with paddy straw produced best results with 95% biological efficiency and high protein content. However, drying of slurry in this case is a problem.
- With the use of detoxified Mahua cake (20%), BGS further increased the quantity and nutritional quality of the mushrooms produced.

# Agaricus:

The compost using BGS produced better quantity and quality (24 Kg / 100 Kg of substrate as compared to 22Kg / 100 Kg of substrate) of mushrooms.

# **BGS As co-composting substrate:**

➤The use of BGS along with the microbial cultures and earthworms reduced the composting time by 35-45% (as compared to the use of BGS alone) and produced enriched compost



# **Future work:**

- 1. Development of Liquid and Solid Biofertilizers i.e. *Rhizobium, Azotobacter, Azospirilum,* Ectomycorrhiza, etc.
- 2. Development of Liquid and Solid Biopesticides (*Trichoderma, Pseudomonas, Bacillus, Paecilomyces,*) using BGS.
- 3.Development of field level **Centrifuge** (giving best results with optimized Parameters) on decreased cost for BGS dehydration purpose.
- 4. More work on Use of Moringa as Coagulant.
- 5. Validation of reports on other uses of BGS.

# THANKS

# Industries / NGOs involved in BGS dehydration by centrifugation

- SKG Sangha : Non profit voluntary organization (150 L)
- KBK Chem Engineering Limited, Pune Maharastra (500 L)
- Torftech Group, Mumbai, Maharastra (500 L)

### **Research Organization**

- Tamil Nadu Agricultural University (400 L)
- Anna University, Chennai (200L)
- Himachal Pradesh Agricultural University, Palampur (200 L)