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Emission and performance analysis of a diesel engine burning cashew nut shell oil bio diesel mixed with hexanol

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Abstract

In this study, the effect of doping hexanol into biodiesel which is from neat cashew nut shell biodiesel oil on the emissions and the performance characteristics was studied in a constant speed diesel engine. The main purpose of this work is to reduce various emissions and also to improve the performance of the diesel engine when fueled with blends of hexanol and neat cashew nut shell biodiesel. Cashew nut shell oil is not edible, and hence it can be used as a viable alternative to diesel. Cashew nut shell biodiesel is prepared by conventional transesterification. Hexanol with 99.2% purity was employed as an oxygenated additive. Experimental studies were conducted by fueling diesel as a baseline and by fueling hexanol and neat cashew nut shell biodiesel mixture. A fuel comprising 10% (by volume) of hexanol and 90% (by volume) neat cashew nut shell biodiesel was referred to as CNSBD900H100 and fuel comprising 20% (by volume) of hexanol and 80% (by volume) of neat cashew nut shell biodiesel was referred to as CNSBD800H200. This study also investigated the possibility of using pure biofuel in an unmodified naturally aspirated diesel engine. The outcome of this study showed that adding hexanol at 10% and 20% (by volume) to cashew nut shell biodiesel results in a reduction in emissions. In addition, a significant improvement in brake thermal efficiency and reduction in brake-specific fuel consumptions were achieved. Hence, it could be concluded that hexanol could be a viable and promising additive for improving the drawbacks of biodiesel when it was used to fuel an unmodified diesel engine.

Keywords Hexanol · Biodiesel · Engine performance · Emission

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1 Introduction

There is concern that availability of crude oil and petroleum products may decline in the future and that they will also continue to be expensive. On the other hand, the fuel economy of engines is improving and will continue to progress. However, the massive increase in the number vehicles has led to high fuel demand. Gasoline and diesel fuels may probably become limited and costly in the near future. An alternate fuel must be developed to reduce the forecast gap between the demand and the availability of fuel. Biofuels derived from vegetable oils are found to be the most promising alternative fuel to diesel. Biofuels can be employed in existing diesel engines with minor or no modifications (Mohod et al. 2015). In addition, the use of biofuel can extend the life of diesel engines because it is more lubricating than petroleum diesel fuel (Mohod et al. 2011). However, studying the behavior of biofuel as a neat fuel is an inevitable task for designing the combustion

process of the future's engines. The chemical composition of these biofuels should also be studied in order to use them as a prospective alternate fuel for future application in engines and combustors. The recent scenario suggests that about 88% of the global energy consumption is from fossil fuels (Mohod et al. 2008, 2011). On the other hand, fossil reserves are predicted to become depleted in the future.

Among the biofuels like biogas, bioalcohol, and biodiesel, alcohol seems to be most common. Biogas requires high pressure for its use in automobile, and its leakage can be hazardous (Rakopoulos et al. 2010a). Biodiesel from edible vegetable oil can cause a shortage in the food supply. Non-edible oil sources require large-scale cultivation which can take up the land resources meant for food crops. Alcohols can be obtained by anaerobic fermentation of lignocellulosic biomass which includes agricultural waste biomass (rice straw, cornstalks, and sugarcane bagasse), forestry biomass (wood-pulp, saw-mill, and paper-mill discards) without much reliance on food crops (Rakopoulos et al. 2011). Further energy crops like switchgrass (Panicum virgatum) and elephant grass can produce a high yield of lignocellulosic biomass that can serve as raw materials for alcohol production. Hence, availability of alcohols would not be an issue.

Ethanol and butanol are suitable for gasoline engines and can be used as octane improvers since their low relative reactivity at low temperatures suppresses uncontrolled auto-ignition (Rakopoulos et al. 2010b). While auto-ignition should be avoided in gasoline engines, it is necessary for diesel engines and for modern engines. Thus, new fuels with high reactivity at low temperatures are required to achieve the overall aim of reducing the world's dependency on conventional fossil fuels and reducing greenhouse gas emissions.

Hexanol is an alcohol with six carbon atoms in the parent chain. It is a colorless liquid, slightly soluble in water, but miscible with diethyl ether and ethanol. Two additional straight-chain isomers of 1-hexanol, 2-hexanol, and 3-hexanol exist, both of which differing by the location of the hydroxyl group (Karabektas and Hosoz 2009; Murcak et al. 2013; Campos-Fernandez et al. 2013). Many isomeric alcohols have the formula $C_6H_{13}OH$. It is used in the perfume industry. The present study explores the possibility of using hexanol as a blend component with neat diesel by up to 20% by volume. Blending of oxygenated alcohols like hexanol with biodiesel will populate fuel-rich regions with hydroxyl group radicals that can catalyze unsaturated HC (hydrocarbon) species to be oxidized rather than participate in soot growth reactions (Murcak et al. 2015; Park et al. 2011).

The other possible reason for choosing hexanol as an alcohol is that hexanol is a 6-carbon straight-chain alcohol that has great potential as a blending component with diesel fuel owing to its higher energy density, higher cetane number, better blend stability and less hygroscopic nature than other widely studied lower alcohols like ethanol, methanol, and even butanol (De Poures et al. 2017). The latent heat of vaporization, density, and viscosity of hexanol is also closer to that of diesel fuel compared to other alcohols (De Poures et al. 2017).

This study paves a way on blending hexanol in neat biodiesel derived from cashew nut shell oil to view its effects on emission characteristics. This study employed four fuels namely, neat diesel, neat cashew nut shell biodiesel, fuel comprising 10% (by volume) of hexanol and 90% (by volume) of neat cashew nut shell biodiesel and fuel comprising 20% of hexanol and 80% of neat cashew nut shell biodiesel in a constant speed compression ignition engine. The emission and performance characteristics of all the test fuels were investigated and compared with the baseline diesel at ambient conditions.

2 Experimental materials and methods

2.1 Cashew nut shell oil

The cashew nutshell is about 0.3 cm thickness with a soft honeycomb structure. Cashew nut shell consists of three layers namely the endocarp, epicarp, and mesocarp. The mesocarp consists of natural resin which holds the shell oil. Cashew nut shell contains a dark reddish brown viscous liquid termed cashew nut shell liquid oil. Cashew nuts shells are a possible feedstock for biodiesel production. Oil extracted from the shell coat and the nut is referred to as cashew nut shell oil (CNSO). The shells of nuts from cashews are dried to reduce the moisture content. They are then milled for higher content of oil. Cashew nut shell oil contains 10% of cardol and 90% of anacardic acid (Mohod et al. 2011). The demand of cashew nut shell oil was found to be 10,000 tonnes. The expected growth pace in demand is 7%-8% per annum (Mohod et al. 2008). Fatty acid composition of biodiesel is shown in Table 1.

Table 1 Fatty acid compositions of biodiesel

Fatty acids	Cashew nut shell methyl ester, vol%
Palmitic C16:0	12.5
Stearic C18:0	6.6
Oleic C18:1	28.9
Linoleic C18:2	35.5
Linoleic C18:3	16.5

2.2 Hexanol

Hexanol is a higher organic alcohol with a six-carbon chain and has a structural formula of CH₃(CH₂)₅OH. Hexanol is a renewable feedstock obtained from glucose and lignocellulosic biomass by means of engineered micro-organisms like Clostridium species (De Poures et al. 2017). Hexanol is obtained by anaerobic fermentation of lignocellulosic biomass such as rice straw, corn stalks and sugarcane bagasse, wood-pulp, saw-mill and paper-mill discards without much reliance on food crops. Further energy crops like switch grass (Panicum virgatum) and elephant grass can produce a high yield of lignocellulosic biomass that can serve as raw materials for hexanol production. Hexanol is slightly soluble in water but miscible with diethyl ether and ethanol. Two additional straightchain isomers of 1-hexanol, 2-hexanol, and 3-hexanol exist, both of which differing by the location of the hydroxyl group (De Poures et al. 2017). For this study, analytical grade hexanol containing the maximum limits of impurities of cyclohexanone (1.1 vol%), water (0.3 vol%) and the rest being traces of other foreign alcohols. Table 2 shows the properties of hexanol employed in this study.

2.3 Transesterification process

Cashew nut shell oil was converted into cashew nut shell biodiesel by conventional transesterification at three different stages.

2.3.1 Acid-catalyzed esterification process

A methanolic solution comprised of 95 mL of methyl alcohol and 5 mL of sulfuric acid was added at a molar

Table 2 Properties of hexanol (molecular formula, C_6H_{13} -OH).Source Merck Millipore, India

Properties	Value
Molecular weight, kg/kmol	102.18
C, wt%	70.52
H, wt%	13.70
O, wt%	15.70
Cetane number	23
Self-ignition temperature, °C	285
Density at 15°C, kg/m ³	821.8
Viscosity at 40 °C, mm/s ²	3.92
Lower heating value, MJ/kg	39.10
Latent heat of evaporation, kJ/kg	486
Vapor pressure, mmHg	1
Boiling point, °C	157
Flashpoint, °C	59

ratio of 1:16 to the cashew nut shell oil. This sample was then heated and maintained at a constant temperature of 60 °C for four different time durations (30, 45, 60 and 90 min) under constant stirring with a magnetic stirrer with a hot plate. Based on the result, it was found that heating the mixture at a temperature below 60 °C for 45 min was the optimum operating condition for the acid-catalyzed esterification process.

2.3.2 Alkali catalyzed transesterification process

The transesterification process was continued using sodium hydroxide alkali catalyst. The use of sodium hydroxide was to reduce the free fatty acid content presented in the cashew nut shell oil. In the second stage, the heated cashew nut shell oil was mixed with sodium hydroxide. This process was carried out at different molar ratios and time durations. The optimum conditions for the alkali transesterification process were found to be: oil to methanol molar ratio of 6:1, concentration of sodium hydroxide of 1 wt%, reaction temperature of 60 °C and reaction time of 45 min while stirring. Through this process, biodiesel was produced from the reaction between vegetable oil and an alcohol with the aid of strong alkyl catalyst.

2.3.3 Purification of the oil

The obtained oil from the alkali transesterification process consisted of cashew nut shell esters and methanol. In the purification process, the methanol dissolved in the oil was removed from cashew nut shell esters to improve the yield of esters. During this process, the transesterified oil was heated at 80 °C obtaining a yield of 90% cashew nut shell oil biodiesel. The properties of tested fuels are listed in Table 3.

2.4 Test engine setup

A 4.2 kW single-cylinder, four-stroke air-cooled diesel engine having a compression ratio of 17 was employed in this study. The engine specification is listed in Table 4. The engine exhaust gases such as NO, CO and hydrocarbons (HC) were measured with the aid of an AVL-444 Di gas analyzer and measured at 300 mm from the exhaust valve. A chemiluminescence detector was employed for detecting NO emission. It produces a linear and equi-molar response to nitrogen compounds by using a stainless steel burner to achieve high-temperature combustion of nitrogen containing compounds to form nitric oxide (NO). A flame ionization detector analyzer was used to detect HC emissions detecting ions formed in the combustion of organic compounds in a sample, producing charged molecules that cause electrical conduction between two electrodes. The

Table 3 Properties of tested fuels

Properties	CNSBD	CNSBD900H100	CNSBD800H200	Diesel	Method
Water content, wt%	0.11	0.10	0.10	0.001	ASTM D2709
Density at 18 °C, gm/cc	0.8833	0.8541	0.8682	0.8210	ASTM D4052
Kinematic viscosity at 35 °C, mm ² /s	4.30	4.18	4.12	2.5	ASTM D445
Calorific value, kJ/kg	38,108	38,884	38,912	42,950	ASTM D240
Cetane index (CI)	52	50	48	46	ASTM D976
Flash point, °C	140	120	111	50	ASTM D93
Iodine value, g/100 g-oil sample	65	65	64	_	ASTM D1510
C, wt%	77.2	78.2	75	_	ASTM D5291
H, wt%	11.4	10.8	12	_	ASTM D5291
O, wt%	11.4	11	13	_	-

Table 4 Specification of	
experimental setup made by	
Kirloskar oil engine Ltd.	
(Khadki, Pune, Maharashtra,	
India)	

Specification	Value
Stroke	4
Cylinder	Single
Rated power	4.2 kW
Rated speed	1300 rpm
Bore diameter (D)	87.5 mm
Stroke (L)	110 mm
Compression ratio	17:1

ions are attracted to a collector plate and induce a current upon hitting the plate and the non-dispersive infrared analyzer was used for detecting CO emissions. Each constituent gas in a sample will absorb some infrared at a particular frequency. By shining an infrared beam through a sample cell (containing CO or CO_2) and measuring the amount of infrared absorbed by the sample at the necessary wavelength, a NDIR detector is able to measure the volumetric concentration of CO or CO_2 in the sample. Smoke opacity was measured by AVL-437C smoke meter after reducing the pressure and temperature in the expansion chamber.

2.5 Uncertainty analysis

Uncertainty, the error of the experiment can occur due to the procedure, experiment condition, calibration as well as the environment. Defining uncertainty is vital in verifying the reliability of experiments. The percentage uncertainties of the experiment were calculated using the linearized approximation method and are ± 4.35 . Measuring accuracy and uncertainty of various equipment are listed Tables 5, and Tables 6, 7 and 8 in "Appendix" show the sample calculation of uncertainties percentage for HC, CO and NO_x emissions. Uncertainty = Square root of [(uncertainty of BP)² + (uncertainty of BSEC)² + (uncertainty of BTE)² + (uncertainty of NO_x)² + (uncertainty of CO)² + (uncertainty of HC)² + (Uncertainty of PM)²] =

$$\sqrt{ \left[(0.75)^2 + (1.8)^2 + (1.45)^2 + (0.45)^2 + (1.1)^2 + (2)^2 + (1.9)^2 \right] }$$

= ± 4.35 %

2.6 Test fuels conditions

Fuel properties were tested for all the tested fuels as per ASTM standards. Engine speed was retained at 1300 rpm during the trial. Errors during the measurement of CO, HC, NO_x and smoke emission were found by the procedure recommended by Moffat et al. (1985).

3 Results and discussion

3.1 Carbon monoxide (CO)

Variation in carbon monoxide emissions with a load for CNSBD1000, CNSBD900H100, and CNSBD800H200 is shown in Fig. 1. CO emissions from biofuels are comparatively less than diesel at all brake power. This is as a result of the abundant availability of inbuilt oxygen in hexanol and cashew shell biodiesel blends (Sahin and Aksu 2015). CO emission decreases with increase in hexanol content. By doping 100 and 200 mL of hexanol to CNSBD, 8.1% and 9.3% of CO emissions were lowered, respectively. The reason for lower CO emissions for CNSBD900H100, CNSBD800H200 is because of improved combustion. Hexanol acts as additional oxygen buffer which stores and

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Table 5 Gas analyzer and smoke meter accuracy details	Parameters	Accuracy	Uncertainty, %
	СО	0.01%	± 1.1
	HC	\pm 2 ppm	± 2
	NO _x	\pm 10 ppm	± 0.45
	Smoke	\pm 1% full-scale reading	± 1.9
	BTE	± 0.5	± 1.45
	Brake-specific fuel consumption (BSFC)	\pm 0.05 g/kWh	± 1.8

CO Carbon monoxide, HC Hydrocarbon, NO_x Nitrogen Oxide, BTE Brake Thermal Efficiency



Fig. 1 Variation of CO with load

releases during combustion (De Poures et al. 2017). The other possible reason for the reduction of CO emissions is owing to reduced viscosity of CNSBD900H100, CNSBD800H200. The fuel with lesser viscosity aids a better evaporation of fuel with air and results in improved combustion & lesser CO emissions. This result is in agreement with many other experimental works (Rajesh Kumar and Saravanan 2015; Aydin and Ogut 2017; Sundar and Saravanan 2011).



Variation in hydrocarbon emissions with a load for CNSBD1000, CNSBD900H100, and CNSBD800H200 is shown in Fig. 2. HC emissions from hexanol and biodiesel blends were lower than diesel at all loads. This attributed to the excess oxygen present in the fuel (Yuvarajan and Venkata Ramanan 2016a).

HC emissions increased with load for all the fuels. At higher loads, the mixture was too rich causing incomplete combustion and higher HC emission (Devarajan et al. 2016). HC emissions drop at all brake power by doping with hexanol. Increasing the hexanol content reduces the HC emissions significantly. This was caused by the lower viscosity of CNSBD by doping hexanol (Devarajan et al. 2017). The viscosity of CNSBD900H100 and CNSBD800H200 was reduced by 1.6% and 1.8%, respectively, comparing neat CNSBD. Due to lower viscosity, the mixing rate of fuel with air was drastically improved causing enhanced combustion and lower HC emission (Venkata Ramanan and Yuvarajan 2016). In addition, the vaporization of hexanol was faster which increased the pace of vaporization (De Poures et al. 2017; Sundar and Saravanan 2011).



Fig. 2 Variation of HC with load



Fig. 3 Variation of NO_x with load



Fig. 4 Variation of smoke intensity with load

3.3 Oxides of nitrogen (NOx)

Variation in NO_x emissions with a load for CNSBD1000, CNSBD900H100, and CNSBD800H200 is shown in Fig. 3. NO_x emission of CNSBD, CNSBD900H100 and CNSBD800H200 was found to be higher than that of diesel at all brake power. Higher oxygen content in the alcohol and biodiesel blends was favorable to the combustion process and consequentially increased its temperature (Britto and Martins 2014; Yuvarajan and Venkata Ramanan 2016b).

 NO_x emissions increase with a load for all test fuels. At higher loads, the temperature attained by the combustion chamber will be high causing higher NO_x emissions (Yuvarajan and Venkata Ramanan 2016c). Further, at higher loads, richer fuel mixture is supplied. By doping 10% and 20% of hexanol to CNSBD, the NO_x emissions were reduced by 3.1% and 4.7%. This is because of lower cylinder temperature attained by these blends as a result of the higher latent heat of vaporization and lower calorific value of hexanol in CNSBD. Further, it is also due to reduced viscosity of the hexanol and biodiesel blends.



Fig. 5 Variation of brake thermal efficiency with load

Lower viscosity of CNSBD and hexanol blend aids better atomization of fuel droplets and generates higher dispersion rate leading to improved combustion rate (lower ignition delay) and thereby reducing the temperature in the combustion chamber (Sundar and Saravanan 2011; De Poures et al. 2017).

3.4 Smoke emissions

Variation in smoke emissions with a load for CNSBD1000, CNSBD900H100, and CNSBD800H200 is shown in Fig. 4. Smoke emissions for hexanol and cashew shell biodiesel blends are lower than diesel at all loads owing to inbuilt excess oxygen content in biofuels (Yuvarajan and Venkata Ramanan 2016c). Smoke emission increases with load for all fuels. At higher load, a higher measure of fuel is required to sustain the constant power output leading higher smoke emissions (Li et al. 2015; Yilmaz and Atmanli 2017).

By doping 10% and 20% of hexanol to CNSBD, smoke emission was reduced by 1.1% and 1.4%, respectively. This is caused by oxygen availability in hexanol which enhances the combustion rate and reduce formation of smoke emissions (Sundar and Saravanan 2011; De Poures et al. 2017).

3.5 Brake thermal efficiency (BTE)

Brake thermal efficiency defines the amount of useful energy obtained by burning a fuel. Variation in BTE with a load for CNSBD1000, CNSBD900H100, CNSBD800H200, and diesel is depicted in Fig. 5. The efficiency of diesel fuel was higher than CNSBD, CNSBD900H100, and CNSBD800H200 at all loads. This is as a result of the lower heating value of biodiesel and alcohol blends (De Poures et al. 2017). The efficiency of all fuels increases with load. At higher loads, the amount of fuel supplied to maintain power is more which consecutively results in higher BTE (Yuvarajan et al. 2017).

It was also found from the figure that the BTE for cashew nut shell biodiesel and hexanol blends (CNSBD900H100 and CNSBD800H200) increases with BTE for CNSBD900H100 hexanol content. and CNSBD800H200 was higher than CNSBD by 0.48% and 0.75%, respectively, at full load conditions. The possible reason could be due to the higher energy density of hexanol which enhances the rate of combustion and ends up with higher BTE (Sundar and Saravanan 2011). Moreover, the viscosity of CNSBD900H100 and CNSBD800H200 was lower than CNSBD1000 causing improved atomization and vaporization of fuel with air leading to higher BTE (Rajesh Kumar and Saravanan 2015; Aydin and Ogut 2017; Sundar and Saravanan 2011).



Fig. 6 Variation of brake-specific fuel consumption (BSFC) with load

3.6 Brake-specific fuel consumption

Brake-specific fuel consumption gives the quantity of fuel required to produce a unit of power output. Variation in brake-specific fuel consumption (BSFC) with a load for CNSBD1000, CNSBD900H100, CNSBD800H200 and diesel is shown in Fig. 6. BSFC increases with a load of all the tested fuel. At higher load, the fuel required to maintain the constant steady power output was higher (Yuvarajan and Venkata Ramanan 2016c). BSFC for diesel was lowest among all test fuels owing to its higher calorific value (Yilmaz and Atmanli 2017).

BSFC for CNSBD900H100 and CNSBD800H200 at full load condition was 0.00752 and 0.01484 kg/kWh lower than CNSBD1000. BSFC reduces with an increase in hexanol content. This was due to increase in energy density of fuel mixtures by doping hexanol. In addition, it was also due to a reduction in viscosity of fuel with the addition of hexanol (Sundar and Saravanan 2011; De Poures et al. 2017). Fuel with lower viscosity assists the combustion process as it combines the fuel with air.

4 Conclusion

The impact of adding hexanol to cashew nut shell biodiesel on emission characteristics was measured and compared with petroleum diesel. The following were the observation derived from this experimental work:

- (1) Hexanol was found to be miscible in cashew nut shell biodiesel.
- (2) CO, HC, smoke emissions of hexanol and neat CNSBD blends were lower than diesel at all loads.
- (3) CO emissions for CNSBD1000 drop by 8.1% and 9.3% when blending it with hexanol of 10% and 20% by volume.
- (4) HC emissions were reduced by 1.6% and 1.8% for CNSBD/hexanol blends at all loads.
- (5) NO_x emissions for CNSBD1000 were 4.71% higher than diesel at loads. However, by adding hexanol to CNSBD1000, NO_x emissions were reduced by 3.1% and 4.7% for at all loads.
- (6) Smoke opacity reduced for CNSBD/hexanol blends by 1.1% and 1.4% when compared to neat CNSBD1000.
- (7) BTE was found to increase marginally by adding CNSBD/hexanol blends. However, it is found lower than diesel at all loads.
- (8) BSFC reduces with increase in the percentage of hexanol in CNSBD/hexanol blends.

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Appendix 1

See Table 6.

			$(HC_{BD}-HC_{D})/HC_{D}$ for different fuels					
	CNSBD1000, %	CNSBD900H100, %	CNSBD800H200, %	CNSBD700H300, %				
0	- 28.85	- 42.31	- 17.31	- 28.94				
25	- 23.75	- 40.00	- 31.25	- 36.75				
50	- 36.67	- 50.00	- 40.00	- 43.67				
75	- 28.57	- 48.57	- 34.29	- 42.86				
100	- 41.67	- 58.33	- 36.25	- 47.50				
Average	- 31.90	- 47.84	- 31.82	- 39.94				

HCBD HC emission of biodiesel-hexanol blends, HCD HC emission of diesel

Table 6Average calculation ofHC emission

Appendix 2

See Table 7.

Table 7Average calculation ofCO emission

Engine load, %	(CO _{BD} -CO _D)/CO _D for different fuels					
	CNSBD1000, %	CNSBD900H100, %	CNSBD800H200, %	CNSBD700H300, %		
0	- 10.82	- 19.52	- 6.14	- 8.15		
25	- 5.82	- 17.20	- 4.59	- 13.78		
50	- 25.18	- 36.03	- 13.54	- 20.16		
75	- 18.89	- 31.30	- 6.52	- 15.96		
100	- 19.33	- 61.73	- 6.67	- 40.00		
Average	- 16.01	- 33.16	- 7.49	- 19.61		

COBD CO emission of biodiesel-hexanol blends, COD CO emission of diesel

Appendix 3

See Table 8.

Fable 8 Sample calculation of uncertainty (%)	NO_x emissions in three tests, ppm			Average NO _x emissions (\pm 1), ppm	Uncertainty, %
	Test 1	Test 2	Test 3		
	363	365	363	364	0.55
	523	521	519	521	0.58
	689	692	693	691	0.43
	808	808	812	810	0.37
	895	899	895	896	0.33
	Uncertaint	y level, %			0.45

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