Status on outcome and utilization of R&D Projects by CSB R & D institut	es in the area of Host plant improvement executed	during the last	10 years (2010-20)
	1 I	0	

#	Project Code and title P	Project	Objectives	Achievement / output	Utilization status of the Coverage / impact		Remarks
	p	period			output	in the field	
1	PIP- 4678- Morpho- physio- anatomical characterization of <i>terminalia spp</i> .	2008 - 2012	1.To characterize the gene pool of <i>Terminalia</i> spp. 2.To isolate superior genotypes of <i>T. arjuna</i> and <i>T. tomentosa</i> from the gene pool for developing superior hybrids.	Isolated 09 superior accessions of <i>T. arjuna</i> .	01 accession of <i>T.arjuna</i> (102) is under multiplication trial and remaining 08 accessions have been included in a Project on "Development of superior hybrids of <i>T.</i> <i>arjuna</i> and <i>T.tomentosa</i> for high leaf yield and quality".	02 saplings each of two improved accessions of <i>Terminalia arjuna</i> viz., 102 and 123 will be supplied to 5 RSRSs & 3 RECS for multiplication. These plants will be compared with the hybrid plants developed under the project PIB-4697 in the proposed project "Evaluation of selected hybrids of <i>Terminalia arjuna</i> and <i>T. tomentosa</i> and drought tolerant <i>T. arjuna</i> accessions, and popularization of superior <i>T. arjuna</i> accessions (acc. 102 & acc. 123) and <i>Lagerstroemia</i> speciosa in different tasar silkworm "ant ragione"	
2	PIB-4686- Studies on 2 various provenances of <i>Terminalia arjuna</i> Bedd. and <i>T.tomentosa</i> W. & A.	2008-2011	1. To study the variability in seeds and growth performance of seedlings of different provenances to estimate the variability among plus trees of T. arjuna and T. tomentosa for	Twenty five plus trees of <i>T. arjuna</i> (11) and <i>T. tomentosa</i> (14) were selected from different locations in Jharkhand and M.P. and evaluated for seed morphology, germination parameters, seedling growth, variability and biomass production. Based on the date 03 plus trees were selected [01 of <i>T.arjuna</i> from Chakradaharpur; Jharkhand 02 of <i>T.tomentosa</i> Bahragoda (Jharkhand) and Mandla (M.P.)	03 plus trees were selected [01 of <i>T.arjuna</i> from Chakradaharpur; Jharkhand 02 of <i>T.tomentosa</i> Bahragoda (Jharkhand) and Mandla (M.P.)]	The information generated is being used to raise uniform population from plus trees.	

			identifying suitable ones for further			
3	PIB-4697 - Development of superior hybrids of <i>Terminalia arjuna</i> and <i>T.</i> <i>tomentosa</i> for high leaf yield and quality.	2012 - 2018	<ol> <li>Identification of most suitable mother plant for raising a seedling orchard with uniform population.</li> <li>Evolution of superior hybrids which can be multiplied vegetatively.</li> </ol>	Superior hybrids identified under pot condition will be evaluated in field trial at different agro- climatic conditions. Based on their performance across the location stable hybrid (s) would be recommended for farmers cultivation.	Identified true hybrids with higher yield traits can be used for mass vegetative multiplication at farmers' level. Project on "Evaluation of selected hybrids of <i>Terminalia arjuna</i> and <i>T.</i> <i>tomentosa</i> and drought tolerant <i>T. arjuna</i> accessions, and popularization of superior <i>T. arjuna</i> accessions (acc. 102 & acc. 123) and <i>Lagerstroemia speciosa</i> in different tasar silkworm rearing regions" has been proposed.	
4	PPA-4704- Development of package for cultivation of <i>Lagerstroemia</i> speciosa for rearing of Tasar silkworm, Antheraea mylitta D.	2014 - 2016	<ol> <li>Development of package for economic cultivation of <i>Lagerstroemia</i> <i>speciosa</i>.</li> <li>Standardization of package for Tasar silkworm (<i>Antheraea</i> <i>mylitta</i> D.) rearing on <i>Lagerstroemia</i> <i>speciosa</i>.</li> </ol>	Propagation through cuttings showed that days taken for rooting varies in different season and maximum days taken for rooting was 45 days in <i>L. speciosa</i> and 60 days in <i>T.arjuna</i> in the cuttings planted during December. Height of one year old saplings of <i>L. speciosa</i> was more (80.13 cm) in comparison to <i>T. arjuna</i> (59.10 cm). Overall growth performance along with total leaf yield/plant of two year old plants of <i>L. speciosa</i> was higher than <i>T. arjuna</i> under 10' x 5' spacing. First crop silkworm rearing results show that larval duration was slightly higher in <i>L.speciosa</i> (35 days) as compared to <i>T.arjuna</i> and <i>T.tomentosa</i> (32 days). Similar trend was observed in second crop of both the year (52 and 45 days in <i>L.speciosa</i> and <i>T.arjuna</i> , respectively). ERR (%) was almost at par in all the three plants in both the crops during 2014 and 2015. Silk ratio was in the range of 11.71 in <i>T.arjuna</i> 13.11 in <i>L.speciosa</i> in first crop during 2014 and 2015. However, silk ratio was higher in second crop during both years in all the three species studied. Filament	Findings revealed that plantation of <i>L.speciosa</i> can be taken up for tropical Tasar culture as additional tropical Tasar silkworm food plant species under block as well as forest plantation. It is very easy to propagate and fast growing as compared to <i>T.arjuna</i> and <i>T.tomentosa</i> . Furthermore, it has an added advantage over <i>T.arjuna</i> that two consecutive rearing can be taken up on same plants. The package of cultivation and silkworm rearing for <i>Lagerstroemia speciosa</i> is being utilized for field recommendation.	

			length varied 630.40 m in <i>T.arjuna</i> to 695m in <i>L.speciosa</i> in first crop of year 2014.Filament length was higher in second crop of year 2015 (887 and 940m in <i>T.tomentosa</i> and <i>L.speciosa</i> , respectively). Denier of the filament was almost similar in all the three species. Grainage behavior in <i>L.speciosa</i> was at par with <i>T.arjuna</i> .			
5	PIC-4705- Development of in-situ soil health and nutrient management in tasar growing areas.	1) To make the farmers aware about the importance of soil fertility for the production of quality leaves and subsequently cocoons through soil health cards.	A farmer friendly package for in-situ soil health and nutrient management developed.	Package is utilized at farmers' field and the same will be adopted by more and more farmers in near future.	Field trials were conducted during 2019-20 at RSRS Dumka, REC CKP and REC Seoni- Champa.	
6	AIP 4711 - Screening of fast growing drought tolerant accessions of <i>T</i> . <i>arjuna</i> for raising block plantation.	1) To select the fast growing accessions of <i>Terminalia arjuna</i> available in the gene bank of CTR&TI for drought tolerance at the early stage of its growth.	Accessions of <i>Terminalia arjuna</i> plant accession no. 525, 523, 123 and 135 could be used as the drought tolerant accessions under limited or severe water stress conditions.	These identified accessions under this project could be useful among the farmers to successful rearing of tasar silkworm and enhance their productivity in the drought prone regions. Project on "Evaluation of selected hybrids of <i>Terminalia arjuna</i> and <i>T.</i> <i>tomentosa</i> and drought tolerant <i>T. arjuna</i> accessions, and popularization of superior <i>T. arjuna</i> accessions (acc. 102 & acc. 123) and <i>Lagerstroemia speciosa</i> in different tasar silkworm rearing regions" has been proposed.		
7	PPS 4725 - Soil Health Cards for Sericulture Farmers.	9 1) To make the farmers aware about the importance of soil fertility for the production of quality leaves and	A combination of secondary nutrients $SM_5$ has been developed and found to increase leaf yield by 27.45% with simultaneous increase in cocoon characters.	Packagefortheapplicationofthesecondarynutrientscombination $SM_5$ forenhancingtheleafyieldandcocooncharacters		

			subsequently cocoons through soil health cards.		was prepared and distributed to all Regional Tasar Research Stations for multi location trial. Large scale field trial at farmer's level is also being taken in Hatgamharia, Jharkhand.	
8	PIT-4700 -Multiplication Jan, of elite accessions of Dec <i>Terminalia arjuna</i> through tissue culture.	n, 2013- ec, 2013	<ol> <li>Standardization and optimization of in vitro technique for micro-propagation of the selected accession / plus trees.</li> <li>In vitro screening of high responding accession / plus trees.</li> </ol>	Standardized the protocol for micro propagation and hardening of elite strain (BLRD) of <i>T. arjuna</i> .	In totality this project has developed the integrated method of elite plant multiplication technique called" In-vitro cum in- vivo juvenile multiplication techniques for tasar host plant "which has a wide application	
9	<b>ARP</b> - <b>4714</b> - 2011 Identification of early sprouting and fast growing genotypes of <u><i>Quercus serrata</i></u> for raising block plantation in North – West India.	16-2020	<ol> <li>Identification of early sprouting and fast growing genotypes of <i>Q.</i> <i>serrata</i> in the existing population.</li> <li>Multiplication of isolated early sprouting genotypes to raise block plantation for utilization in early spring crop (March – April).</li> </ol>	i) Five plants were identified as early sprouters during four surveys conducted under the project. Out of total 5 identified plants, three were identified from Kumaon and two from Garhwal region. No rooting was observed in air layers tried on the plants. Twigs of selected genotypes were brought to RTRS Bhimtal and planted by appropriate methods. But even after repeated attempts and following different protocols, rooting was not observed. The PI also consulted the Horticulture Dept. G.B. Pant University, Pantnagar and Department of Plant breeding & Tree development, FRI, Dehradun. As suggested, treatment of 4000 ppm IBA with talcum powder & ethyl alcohol before sprouting was also tried but no positive result was found.	During the Institute review meeting on 04.09.2020, it was suggested to try the multiplication of identified early sprouters through seedlings under Programme of work and identify more such plants and try for their multiplication.	

Status on outcome and utilization of R&D Projects by CSB R & D institutes in the area of Silkworm improvement executed during the last 10 years (2010-20)

<b>#</b> Project Code and title		Project	Objectives	Achievement / output	Utilization status of the	Utilization status of the Coverage / R		
		period			output	impact in the		
		-			_	field		
1	AIG-4669- Selection	2006-2011	1. To find out	• Cocoon and shell weight specific	The identified shell	Basic		
	aided molecular marker		association of molecular	markers have been identified which are highly	weight linked SCAR	information		
	system for improvement		marker (s) with yield	reproducible and can be used with grater	markers are being used in	generated		
	in tropical tasar		traits of tasar silkworm	fidelity for inheritance pattern analysis of	marker assisted selection	C		
	silkworm Antheraea		2. To assess DNA	vield traits. This is the first report of	for cocoon yield			
	<i>mylitta</i> Drury.		polymorphism and	development of SCAR marker, diagnostic to	improvement in tasar			
	, , , , , , , , , , , , , , , , , , ,		heterozygosity found at	specific vield parameters of A. mvlitta. The	silkworm.			
			individual and	SCAR marker SCOPW-16 <sub>826</sub> could be				
			population level.	effectively utilized to circumvent the problem				
				of lab-to-lab reproducibility and dominant				
				nature of inheritance in RAPD. The				
				discrimination between HCSW and LCSW				
				trait group achieved by these markers would				
				make them as very useful diagnostic markers				
				for silk yield improvement, breeding and				
				marker assisted selection aiming at the				
				development of A. mylitta for productivity.				
				The differentially expressed mRNA patterns				
				in tasar silkworm larvae are very interesting,				
				because some of them are unique to this				
				species				
2	ARP-4793- Studies on	2012 - 2015	1. To study the life	• The grainage technology for wild ecorace	Outcome of the project is	Laria		
	the biology and ecology		cycle, ecology, ethology	Laria have been standardized through which	being utilized for Laria	Conservation		
	of Laria ecorace of		and population dynamics	Laria dfls can be produced.	conservation.	model developed		
	Antheraea mylitta D. on		of Laria ecorace in	• A package of practice for the efficient		and provided to		
	Sal flora		natural habitat.	utilization of vast Sal flora based on the		DOS Jharkhand		
				studies of Laria ecorace biology and		who		
			2. To standardize the	ecology in situ and ex situ is suggested.		implemented the		
			rearing schedule of Laria	<ul> <li>Through phase-wise brushing, it is observed</li> </ul>		same and got 30-		
			ecorace for utilization of	that if brushing is conducted during Rainy		35% success in		
	Sal flora		season (July), there will be less success in	success in conserv				
				productivity.				
				• During autumn season, the successful				
				brushing period is third to fourth week of				
				September. This can be recommended for				
				the area where Laria is being conserved.				
				Cocoons produced during first season help				

## 1. <u>CTR&TI, Ranchi</u>

				<ul> <li>in selective conscription of progenies in the next season with reasonably good survivability if they are left as such untouched and undisturbed by human beings.</li> <li>During Rainy season, Laria should be allowed to proliferate in natural pockets. <i>Ex situ</i> rearing is not much successful.</li> <li>High phenols, tannin content in leaves may be among the causative factors for low gut enzyme activity leading to low protein accumulation. The exact enzyme inhibitor(s) need further studies.</li> <li>Emphasis must be given for conservation of Laria in natural eco-pockets in forests of Jharkhand.</li> </ul>			
3	AIB-4694- Improvement of Daba ecorace of <i>Antheraea</i> <i>mylitta</i> Drury for higher fecundity	2012 - 2015	To improve the semi- domestic Daba race for higher fecundity in tropical tasar silkworm <i>Antheraea mylitta</i> Drury	<ul> <li>Three breeding lines with &gt;230, &gt; 250 and &gt;270 were isolated/developed without detrimental effect to male shell weight and female pupal weight. Developed lines are having higher fecundity when compared with ruling Daba fecundity (200-220).</li> <li>Multi-location trial of breeding line (&gt; 250 eggs/dfls) named as CTR-14 were carried out in different agro-climatic conditions namely RTRS Baripada (Odisha), REC-Katghora (Chhattisgarh), REC-Purulia (West Bengal) and REC-Hatgamaria (Jharkhand) in both the crops.</li> <li>It is recommended that developed lines may be multiplied at P4 station of CTR&amp;TI, Ranchi and good performers are required to be introduced into seed channel through P4 for improving fecundity and in turn cocoon production at farmer's level.</li> <li>A package of practice for rearing of high fecundity line CTR-14 was also worked out and recommended.</li> </ul>	Multiplication trials of improved tasar silkworm line, CTR-14 for higher fecundity are being conducted at RSRS, Dumka, Baripada, Jagdalpur, Warangal & Bhandara and REC, Kathgora (Siwani), Robertsganj Kapistha & Hatgamharia/ Chakradharpur.	As per the suggestions of Hybrid Authorization Committee (HAC), project for validation trial of CTR-14 has been submitted.	
4	AIP-4696- Management of abiotic factors to regulate emergence in dianausing seed	2012- 2015	1) To find a mechanism to check erratic emergence and regulate moth emergence	Temperature treatment at 20oC delayed the emergence.	Temperature treatment at 20°C for 15 days delays the moth emergence in tasar silkworm coccors	The technology is under trial as On Farm Trial (OFT) during	
	cocoons.		with the help of controlled regime of	comparatively found more sensitive to delay in emergence.		2020-21.	

			temperature and humidity from the diapausing seed cocoons	Temperature treatment in month of June is less effective. Except delay in emergence period, no significant variation on emergence (%), fecundity and hatching (%) was observed between control and treated groups.			
5	AIT-4702 - Molecular cloning and heterologous expression of <i>Antheraea mylitta</i> cocoonase.	June, 2013 - May, 2015	<ol> <li>Large scale Collection of cocoonase</li> <li>To express the cocoonase gene in heterologous hosts for its future utilization in softening of tasar cocoons.</li> </ol>	Large scale rearing of <i>A. mylitta</i> and 2150 ml of cocoonase collection from 3000 emerging tasar silkworm has been done successfully. Prior to present study cocoonase was the unutilized by-product of tasar industry. Molecular cloning has been done. The fragment released from PGMT-Clones by EcoR1 digestion was gel eluted and sequenced. Expression of cocoonase was evaluated through SDS-PAGE. Native cocoonase secretion process, place of secretion, secretion volume, and sequential changes in cocoonase concentration, chronological secretion profile, quantification, purification, proper storage conditions, enzymatic action and enzyme activity has been worked out successfully. SDS-PAGE analysis of purified cocoonase showed molecular weight 26 kDa.	Standardize the cocoonase collection from emerged moth. Cloning of cocoonase enzymes.	Second phase of the project "Identification most-active cocoonase of sericigenous insects and its variant through molecular characterization" , funded by DBT is under progress. During the project, efforts will be made to develop commercial use of cocoonase enzyme. (Project code AIT 4728)	
6	ARP 4713 - Isolation of Thermo-tolerant line(s) of tasar Silkworm <i>Antheraea mylitta</i> Drury through molecular studies.	2016 - 2018	<ol> <li>To isolate thermo-tolerant line(s) of tropical tasar silkworm, Antheraea mylitta.</li> <li>To develop molecular marker for the identification of thermo- tolerant line(s).</li> <li>To unravel the underlying molecular mechanism of thermal stress tolerance in tasar silkworm.</li> </ol>	Developed SCAR Markers would be an useful tool for identifying thermo-tolerant lines. Understanding of molecular mechanism behind the thermal tolerance would give an insight for the basis of adaptation in tasar silkworm. The 2 <sup>nd</sup> phase project has been approved by CSB and given project code [AIT04002SI] (February, 2019- January 2022) and has also been initiated.	The isolated thermo- tolerant lines could be utilized successfully in hotter zones of India to overcome the loss of dfl production due to high temperature.	Second phase of the Project AIT04002SI is under progress	
7	AIB 4717 - Improvement of tropical tasar silkworm for high	Oct, 2016 - Sept, 2019	1) Improvement of tasar silkworm breeds for high silk yield	Analysis of variance in RBD revealed significant variation across crosses for male cocoon weight and pupa weight and not for	The finding of the project has been utilized in formulation of a new	A long duration breeding programme has	

	silk yield through recurrent selection.		through recurrent selection breeding.	remaining. This indicates the breeding material has attained genetically homogeneous. Therefore creation of genetic variation in breeding population is essential for further genetic improvement of population. To achieve this evaluated population needs to be crossed with Wild Daba or other cultivated Daba in next generation. However, mean performance of total crosses of second season was found to be better as compared to that of first crop crosses for all the traits.	project proposal entitled "Rejuvenation programme of ruling Daba using wild Daba in Antheraea mylittta its homeland in Singhbhum Jharkhand".	to be taken for development of a high yielding breed which can be successful in the field.	
9	AIB 4687 Improvement of Andhra ecorace of <i>Antheraea</i> <i>mylitta</i> D through back cross method- RTRS, WARANGAL	2008 - 2012	1. To introgress the survival characters of Daba TV ecorace in Andhra ecorace having high superior commercial characters 2.To develop breeds best suited for on farm conditions	Evolved BC-IV line of Andhra Local which yielded 35 cocoons/dfl over the parents (Andhra Local – 12 cocoons and Daba TV – 20 cocoons/dfl).	Increase in cocoon production from 12 (Andhra Local) to 35 (improved line) has been achieved.		

SL	<b>Project code</b>	Project Title	Project	Outcome	Present status
No.			period		
1		Screening, Characterization and Identification of Disease Tolerant Variety in Tropical Tasar Silkworm of <i>A. mylitta D.</i>	2003 - 2006	<ul> <li>Based on repeated observations and LC50, ecoraces grouped as, Tolerant: Modal,Sukinda, Daba BV; Moderate: Sarihan, Daba TV, Bhandara, Andhra local; Susceptible: Raily.</li> <li>The healthy stock (Tolerant) indicates specific bands at the RAPD fragment sizes of 500 bp for Daba, 750 bp for Sarihan, 1 kb for Sukinda and for Andhra ecorace which were not present in corresponding diseased (Susceptible) stock of the ecorace.</li> <li>Polymorphic bands showing specific variation in RAPD fragment sizes ranging between 3kb to 2.5 kb and 2 kb to 2.5 kb between tolerant and suscentible line.</li> </ul>	<ul> <li>Ecorace specific RAPD markers had been recognized for the identification of eight <i>A. mylitta</i>ecoraces.</li> <li>AmCPV tolerance in the different ecoraces found that Modal&gt;Sukinda&gt;DabaBV&gt;DabaTV&gt; Sarihan&gt;Raily and information would be utilized in the development of AmCPV tolerant breeds.</li> </ul>
2		Phylogeography of Antheraeamylitta (Tropical tasar silkworm) and Antheraeaassamensis (Muga silkworm)	2005 – 2008	<ul> <li>Molecular studies indicated significant genetic variability among and within seven ecoraces.</li> <li>Seven ecotypes of the Indian tasar silkworm, <i>A. mylitta</i> analysed using ISSR and SSR primers showed inter- and intra population polymorphism. Daba reared at CTR&amp;TI Ranchi and Daba natural collected from West Singhbhumshowed close proximity indicating domestication did not cause significant genetic variability. This is supported by similar mating behaviour by the semi-domestic and wild populations. Among the wild populations, Modia and Sarihan showed genetic closeness</li> </ul>	• Acquired information is helpful in conservation of wild ecoraces with a conservation genetic approach. Laria conservation programme is based on this information

## Status on outcome and utilization of R&D Projects by CSB R & D institutes in the area of Biotechnology executed during the last 10 years (2010-20)

				while Laria, JataDaba and Bogai clustered to form a distant genetic group	
3	AIG 4669	Selection aided molecular marker system for improvement in Tropical tasar silkworm <i>A. mylitta Drury</i>	2007-2010	<ul> <li>form a distant genetic group.</li> <li>Cocoon weight and shell weight specific markers have been identified which are highly reproducible and can be used with greater fidelity for inheritance pattern amongst yield traits. This SCAR marker, diagnostic to specific yield traits of <i>A.mylitta</i>.could be effectively utilized to circumvent the reproducibility and dominant nature of inheritance in RAPD.</li> <li>Discrimination between high and low cocoon and shell weight group achieved by these markers make them useful diagnostic markers for silk yield improvement, Marker Assisted Selection aiming at the development of <i>A. mylitta</i> productivity</li> <li>The differentially expressed m-RNA patterns in tasar silkworm larvae are very interesting because some of these are unique to this species. This has a relationship with adaptive significance and life cycle strategies of tasar</li> </ul>	The SCAR marker associated with cocoon and shell weight could be utilized for Marker Assisted Selection programme.
4	AIP-4680	Ecogenetic analysis of diapause and reproduction in tropical tasar silkworm <i>A. mylitta Drury</i>	2007-2010	<ul> <li>Slikworm.</li> <li>The role of physical and environmental and genetic factors responsible for pupal diapause and mechanism was studied.</li> <li>Role of bioactive molecules secreted by the male accessory gland responsible for enhancing fecundity was explored.</li> <li>23 EST sequences submitted to NCBI Database.</li> <li>In order to identify the physical and environmental and genetic factors responsible</li> </ul>	Project was exploratory in nature.

5	ARP 4681	Identification and characterization	2007-2010	for pupal diapause and mechanism in A. mylitta, the larval growth and development pattern of the Dababivoltineecorace of A. mylitta was studied on the basis of instar specific green & dry weight, green & dry weight gain in non-diapause-destined (NDD) and diapause-destined (DD) generations along with available photoperiod/day length, relative humidity and temperature, it was found that the green and dry weight gain was higher during early instars and significantly higher growth was observed in the first half of each instar. There was no significant sex-specific variation of green and dry weight of larvae of the same generation.	• The amplicon obtained in 16s rPNA
		of disease causing pathogens (microspordia and bacterial infections) of tropical tasar silkworm, <i>Antheraeamylitta D</i>	2007-2010	• Micosporidia and 19 bacteria isolated from tasar silkworm collected from different geographical tasar growing areas. The mirospordia and bacteria characterized morphologically and biochemically have also shown variation with molecular characterization (RAPD and 16s rRNA gene). The amplicon obtained in 16s rRNA analysis were sequenced and obtained sequences deposited to NCBI database with different accession numbers.	• The amplicon obtained in 16s rRNA analysis were sequenced and obtained sequences deposited to NCBI database with different accession numbers.
6	AIT-4702	Molecular cloning and heterologous expression of <i>A. mylitta</i> cocoonase.	2013 -2015	• Large scale rearing of <i>A. mylitta</i> and 2150 ml of cocoonase collection from 3000 emerging tasar silkworm has been done successfully. Prior to present study cocoonase was the unutilized by-product of tasarindustry.Identification of crucial stage for cocoonase collection was done based on temporal changes in colour of pupae	• Phase-I completed successfully and second phase DBT funded project AIT 4727 (2018-2021) initiated as identification of most active cocoonase for its eventual commercial use. For prediction of protein structure from the sequence procured was constructed from the

				integument. Native cocoonase secretion process, place of secretion secretion volume, sequential changes in cocoonase concentration,	newly constructed sequence. The blast result obtained for particular sequence was subjected for contig
				quantification, purification by sephadexG100 column, characterization, proper storage	baser. In silico analysis of AmCoc gene was done. The structure of
				activity has been worked out successfully. SDS-PAGE analysis of purified cocoonase showed molecular weight 26 kDa. Per insect	Quark (ab initio protein folding and protein structure prediction) server determines 3D structure of protein.
				cocoonase collection volume is 500 to 800µl volume with 221µg/ml concentration. Recognition of sericin as natural substrate of	The TM- score for the best model was observed as $(0.3461 \pm 0.0833)$ .
				cocoonase and it is established that cocoonase directly acts on the sericin without affecting the fibroin protein. SEM study showed marked	
				softening was done using cocoonase. MaldiTof-Tof (MS and MSMS) data A& B of A mylittacocoonase MS 1320 477 showed	
				similarities with cocoonase/proteolytic enzyme of other sericigenous insects. Complete coding nucleotide sequences of cocoonase gene were	
				retrieved from NCBI database and phylogenetic analysis conducted. Sequence annotation studied was conducted and	
				sequence was submitted to NCBI (>gi 731516038 gb KM388539.1  UNVERIFIED: Antheraeamylitta genomic	
				sequence). Molecular matching of Antheraea spp. cocoonase was also performed in order to identify the eventual analogues available in	
7	ARP 4713	Isolation of thermo-tolerant line(s)	2016-18	A state of the state of th	• Thermo-tolerant lines could be

		of to got all my and Authouse a multitude		horrs hoon colorial through high the second stress	atilized an each fully in hotton
		of tasar sılkworm <i>Antheraeamylitta</i> <i>Drury</i> through molecular studies		<ul> <li>have been selected through high thermal stress.</li> <li>Three SCAR markers (TT-PB1, TT-PB2 and TT-PB3) have been developed. Among three, TT-PB1 showed more specificity towards thermo-tolerant lines.</li> <li>Heat shock proteins and factors responsible for thermotolerance in thermo-tolerant line have been analysed. HSP70 and HSP21 have found to be upregulated in thermo-tolerant lines of Daba.</li> <li>Expression pattern of α&amp;β-esterase isozyme in fifth instar temperature treated and untreated male and female larval hemolymph of A. mylitta has been analysed. Esterase at Rf value 0.48 has found to be expressed only in temperature treated larvae.</li> </ul>	<ul> <li>utilized successfully in hotter zones of India to overcome the loss of dfl production due to high temperature.</li> <li>Developed SCAR Markers would be a useful tool for identifying thermo- tolerant lines.</li> <li>Understanding of molecular mechanism behind the thermal tolerance would give an insight for the basis of adaptation in tasar silkworm.</li> </ul>
8	AIT-4724	Isolation and characterization of sericin from tasar silk waste for Commercial utilization.	2016-2018	<ul> <li>Boiling with 0.2% sodium carbonate removes the residual sericin. Residual sericin was 2.33- 2.79%. Fibre waste sericin has anti-tyrosinase, anti-elaste and antioxidant properties.</li> <li>Higher quantity of sericin has been isolated from Dabaecorace. Elemental compositions of different ecorace-specific sericin are 11.94- 12.39% Nitrogen, 40.3-40.75% Carbon, 0.715- 1.47% Sulphur and 6.33-6.55% Hydrogen. No significant variation of molecular weight distribution of protein was observed between different tasarecoraces. Railyecoracesericin found to posses more</li> <li>non-polar and aromatic amino acids. Railysericinposses higher inhibition percentage of tyrosinase, glutathione- stransfease activity and hydrogen peroxide</li> </ul>	• For further utilisation of project findings Phase-II of project entitled "II phase project: Mass level extraction of sericin from tasar cocoon cooking waste water for its prospective utilization- sent to DBT New Delhi for funding.

				scavenging notential	
				• Higher phenolic and C: N ratio was observed	
				in sericin separated from	
				• cocoons of Sal food plant. No significant	
				variation of molecular weight distribution of	
				protein and thermal stability was observed in	
				sericin separated from cocoons of different	
				plants. More non-polar amino acids were	
				• observed in case of Sal sericin. As compared	
				with other food plants, serie separated from	
				cocoons of Sal fed larvae posses higher	
				scavenging potential.	
				• The waste water sericin having higher amount	
				of low molecular weight proteins (>35kDa)	
				and lower amount of high molecular weight	
				(<245kDa). Separated sericin was confirmed	
				ETIR spectra and	
				• FLISA test Higher anti-tyrosinase activity	
				DPPH scavenging potential and inhibition of	
				lipid peroxidation potential was also detected	
				in tasar waste sericin.	
9	PIG-4682	Evaluation of gene pool of tropical	2007-2010	<ul> <li>Morphological and biochemical</li> </ul>	• High level of heterozygosity was
		Tasar silkworm host plants with		characterization	found in both the species. Based upon
		Toger gilk		• Analysis of variance for different phenotypic	overall characterization and
		Tasar siik.		and biochemical traits showed significant	evaluation, four promising accessions
				differences among accessions of both 1. arjuna	multiplication and popularization
				among the accessions.	mentproduton and popularization.
				• Cluster analysis indicated that geographical	
				distribution and genetic divergence did not	
				follow the same pattern in both the species.	

<ul> <li>a geographical locations into one cluster appears to be due to presence of some common genes controlling the most important characters through modifying effect of micro- and macro- environment, while the geotypes from same location were grouped in separate cluster indicating occurrence of wide diversity among genotypes from sAme location.</li> <li>Molecular characterization RAPD profiling of 18 accessions of T.arjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic (91.89%) and 52 bands were polymorphic (91.89%) and 52 bands were polymorphic (91.89%) and 52 bands were generated a total of 641 RAPD bands. Among total bands, 589 bands were generated per primers ranging between 7-30 bands. Only 10 primers (OPM-043, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were golymorphic (96.38%) and 26 bands were golymorphic (96.38%) and 26 bands were golymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primers (OPM-03, OPM-07, OPM-06, OPM-11, OPM-13, OPM-14, OPM-17, OPM-06, OPM-11, OPM-11, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-14, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-13, OPM-14, OPM-17, OPM-109, OPM-11, OPM-14, OPM-17, OPM-143, OPM-143, OPM-17, OPM-109, OPM-11, OPM-143, OPM-143,</li></ul>		The grouping of genotypes of different	
<ul> <li>b be due to presence of some common genes controlling the most important characters through modifying effect of micro- and macro- environment, while the genotypes from same location were grouped in separate cluster indicating occurrence of wide diversity among genotypes from sAme location.</li> <li>• Molecular characterization RAPD profiling of 18 accessions of T arjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic</li> <li>(91.89%) and 52 bands were polymorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7.30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPM-11, OPM-16, OPW- 08, OPW-12, OPM-11, OPM-16, OPW- 08, OPW-12, OPM-13, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T. tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05, Among 37 primers only 20 primers (OPM-03, OPM-04, OPM-04, OPM-11, OPM-13, OPM-14, OPM-04, OPM-04, OPM-11, OPM-13, OPM-14, OPM-04, OPM-04, OPM-11, OPM-13, OPM-14, OPM-04, OPM-04, OPM-11, OPM-13, OPM-14, OPM-04, OPM-04, OPM-11, OPM-13, OPM-04, OPM-04, OPM-04, OPM-11, OPM-04, OPM-04, OPM-04, OPM-04, OPM-11, OPM-04, OPM-04, OPM-04, OPM-04, OPM-04, OP</li></ul>		geographical locations into one cluster appears	
<ul> <li>a controlling the most important characters</li> <li>through modifying effect of micro- and macro- environment, while the genotypes from same location were genotypes from same location gocurrence of wide diversity among genotypes from sAame location.</li> <li>• Molecular characterization</li> <li>RAPD profiling of 18 accessions of T.arjunausing 35 random RAPD primers</li> <li>generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic</li> <li>(91.89%) and 32 bands were polymorphic</li> <li>(91.89%) and 32 bands were monomorphic. On an average 16.83 polymorphic bands were generated a total of 641 RAPD bands. Among total bands, 589 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomontosauxing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic. (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated a total of 719 RAPD bands, maximu (35) generated by primer OW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM-17, OPM-09, OPM-11, OPM-10, OPM-01, OPM-02, OPM-10, OPM-11, OPM-13, OPM-01, OPM-02, OPM-10, OPM-11, OPM-10, OPM-01, OPM-02, OPM-10, OPM-10, OPM-11, OPM-13, OPM-01</li> </ul>		to be due to presence of some common genes	
<ul> <li>through modifying effect of micro- and macro- environment, while the genotypes from same location were grouped in separate cluster indicating occurrence of wide diversity among genotypes from sAame location.</li> <li>Molecular characterization</li> <li>RAPD profiling of 18 accessions of T. adjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, S89 bands were polymorphic</li> <li>(91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315)</li> <li>generated 100% polymorphic products The RAPD profiling of 16 accessions of T. tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic (06.38%) and 26 bands were monomorphic 0.01 an average 18,73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer QM-03, OPM-04, OPM-04, 17, OPM-18, OPM-19, OPM-01, OPM-04, 0PM-04, OPM-01, OPM-01, OPM-04, 0PM-04, OPM-11, OPM-13, OPM-04, 0PM-04, OPM-14, OPM-14, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-01, OPW-01, OPM-04, OPM-19, OPW-01, OPW-01, OPW-01, OPM-04, OPM-19, OPM-01, OPW-01, OPM-04, OPM-04, 0PM-04, OPM-19, OPM-01, OPM-01, OPM-04, OPM-04, OPM-19, OPM-01, OPM-01, OPM-01, OPM-01, OPM-04, OPM-19, OPM-01, OPM-01, OPM-01, OPM-04, OPM-19, OPM-01, OPM-</li></ul>		controlling the most important characters	
<ul> <li>antodyning enteror provide the genotypes from same location were grouped in separate cluster indicating occurrence of wide diversity among genotypes from sAame location.</li> <li>• Molecular characterization RAPD profiling of 18 accessions of T. arjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic (91.89%) and 25 bands were polymorphic. On an average 16.83 polymorphic bands were generated a total of 641 RAPD bands. Construction of the construction</li></ul>		through modifying effect of micro- and macro-	
<ul> <li>Indicating courrence of wide diversity among genotypes from sAme location</li> <li>Molecular characterization</li> <li>RAPD profiling of 18 accessions of T. arjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, S89 bands were polymorphic</li> <li>(91.89%) and 52 bands were polymorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30</li> <li>bands. Only 10 primers (OPM-03, OPM-04, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW-08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic primers generated per primer ranging between 7-30</li> <li>bands were polymorphic (96.38%) and 26</li> <li>bands were polymorphic (96.38%) and 26</li> <li>bands were polymorphic (96.38%) and 26</li> <li>bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW-03, OPM-07, OPM-11, OPM-13, OPM-14, OPM-17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPM-11, OPM-13, OPM-14, OPM-17, OPM-18, OPM-11, OPM-13, OPM-14, OPM-17, OPM-18, OPM-11, OPM-13, OPM-14, OPM-15, OPW-10, OPW-1</li></ul>		anvironment while the construct from some	
<ul> <li>Ideation Repaired Brouged in Separate Unsert</li> <li>indicating occurrence of wide diversity among genotypes from sAame location.</li> <li>Molecular characterization</li> <li>RAPD profiling of 18 accessions of</li> <li>T. arjunausing 35 random RAPD primers</li> <li>generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic</li> <li>(91.8%) and 52 bands were polymorphic and by average 16.83 polymorphic bands were</li> <li>generated per primer ranging between 7-30</li> <li>bands. Ohy 10 primers (OPM-03, OPM-04, OPM-04, OPM-05, OPM-07, OPM-10, OPM-16, OPW-08, OPM-04, OPM-05, OPM-07, OPM-10, OPW-08, OPM-04, OPM-05, OPM-07, OPM-10, OPM-16, OPW-08, OPM-04, OPM-06, OPM-06, OPM-06, OPM-06, OPM-06, OPM-06, OPM-07, O</li></ul>		losotion were grouped in concrete eluster	
<ul> <li>Indicating occurrence of whice dressly among genotypes from same location.</li> <li>Molecular characterization</li> <li>RAPD profiling of 18 accessions of Tarjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic</li> <li>(91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW-08, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW-08, OPM-01, OPM-0</li></ul>		in tigeting a segment of soil a discourse	
<ul> <li>Molecular characterization</li> <li>Molecular characterization</li> <li>RAPD profiling of 18 accessions of</li> <li>T.arjunausing 35 random RAPD primers</li> <li>generated a total of 641 RAPD bands. Among</li> <li>total bands, 589 bands were polymorphic</li> <li>(91.89%) and 52 bands were monomorphic. On</li> <li>an average 16.83 polymorphic bands were</li> <li>generated per primer ranging between 7.30</li> <li>bands. Only 10 primers (OPM-03, OPM-04,</li> <li>OPM-05, OPM-07, OPM-11, OPM-16, OPW-</li> <li>08, OPW-12, OPW-18, and AM773315)</li> <li>generated 100% polymorphic products</li> <li>The RAPD profiling of 16 accessions of</li> <li>T. tomentosausing 37 random RAPD primers</li> <li>generated a total of 719 RAPD bands of which</li> <li>693 bands were polymorphic. On an average 18.73</li> <li>polymorphic bands were generated per primer</li> <li>ranging between 11-35 bands, maximum (35)</li> <li>generated by primers (OPM-03, OPM-07, OPM-04, OPM-07, OPM-11, OPM-13, OPM-14, OPM-17, OPM-18, OPM-11, OPM-13, OPM-14, OPM-17, OPM-18, OPM-19, OPM-19, OPM-19, OPM-19, OPM-10, OPW-02, OPW-10, OPW-10, OPW-10, OPW-10, OPW-02, OPW-10, OPW-11, OPW-11, OPW-11, OPW-13, OPM-14, OPM-17, OPM-18, OPM-11, OPM-14, OPM-17, OPM-18, OPM-19, OPM-19, OPM-10, OPW-02, OPW-10, OPW-10, OPW-10, OPW-02, OPW-10, OPW-10,</li></ul>		indicating occurrence of white diversity among	
<ul> <li>Molecular characterization</li> <li>RAPD profiling of 18 accessions of T. arjumausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, S89 bands were polymorphic</li> <li>(91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T. tomentosausing 37 random RAPD primers generated atotal of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-13, OPM-14, OPM- 17, OPM-180, OPM-19, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-13, OPM-14, OPM- 15, OPW-18, OPW-19, AM773371 and</li> </ul>		genotypes from sAame location.	
RAPD profiling of 18 accessions of         T.arjunausing 35 random RAPD primers         generated a total of 641 RAPD bands. Among         total bands, 589 bands were polymorphic         (91.89%) and 52 bands were monomorphic. On         an average 16.83 polymorphic bands were         generated per primer ranging between 7-30         bands. Only 10 primers (OPM-03, OPM-04,         OPM-05, OPM-07, OPM-11, OPM-16, OPW-         08, OPW-12, OPW-18, and AM773315)         generated 100% polymorphic products         The RAPD profiling of 16 accessions of         T.tomentosausing 37 random RAPD primers         generated a total of 719 RAPD bands of which         693 bands were polymorphic (96.38%) and 26         bands were monomorphic. On an average 18.73         polymorphic bands were generated per primer         ranging between 11-35 bands, maximum (35)         generated by primer OPW 05. Among 37         primers only 20 primers (OPM-03, OPM-07,         OPM-09, OPM-11, OPM-13, OPM-14, OPM-17,         OPW-10, OPW-11, OPW-12, OPW-13, OPW-12,         OPW-10, OPW-11, OPW-12, OPW-13, OPW-14,		• Molecular characterization	
Tarjunausing 35 random RAPD primers generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic         (91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-13, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		RAPD profiling of 18 accessions of	
generated a total of 641 RAPD bands. Among total bands, 589 bands were polymorphic (91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		T.arjunausing 35 random RAPD primers	
total bands, 589 bands were polymorphic (91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW-13, OPW-13, OPW-14, OPM- 15, OPW-18, OPW-19, AM773371 and		generated a total of 641 RAPD bands. Among	
<ul> <li>(91.89%) and 52 bands were monomorphic. On an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW-15, OPW-15, OPW-18, OPW-19, AM773371 and</li> </ul>		total bands, 589 bands were polymorphic	
an average 16.83 polymorphic bands were generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T. <i>iomentosausing</i> 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPM-13, OPM-13, OPM- 15, OPW-18, OPW-19, AM773371 and		(91.89%) and 52 bands were monomorphic. On	
generated per primer ranging between 7-30 bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		an average 16.83 polymorphic bands were	
bands. Only 10 primers (OPM-03, OPM-04, OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		generated per primer ranging between 7-30	
OPM-05, OPM-07, OPM-11, OPM-16, OPW- 08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primers OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-107, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		bands. Only 10 primers (OPM-03, OPM-04,	
08, OPW-12, OPW-18, and AM773315) generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		OPM-05, OPM-07, OPM-11, OPM-16, OPW-	
generated 100% polymorphic products The RAPD profiling of 16 accessions of T.tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		08, OPW-12, OPW-18, and AM773315)	
The RAPD profiling of 16 accessions of T. tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-10, O		generated 100% polymorphic products	
T. tomentosausing 37 random RAPD primers generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		The RAPD profiling of 16 accessions of	
generated a total of 719 RAPD bands of which 693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		T.tomentosausing 37 random RAPD primers	
693 bands were polymorphic (96.38%) and 26 bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		generated a total of 719 RAPD bands of which	
bands were monomorphic. On an average 18.73 polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		693 bands were polymorphic (96.38%) and 26	
polymorphic bands were generated per primer ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		bands were monomorphic. On an average 18.73	
ranging between 11-35 bands, maximum (35) generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-01, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		polymorphic bands were generated per primer	
generated by primer OPW 05. Among 37 primers only 20 primers (OPM-03, OPM-07, OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		ranging between 11-35 bands, maximum (35)	
primers only 20 primers (OPM-03, OPM-07,           OPM-09, OPM-11, OPM-13, OPM-14, OPM-           17, OPM-18, OPM-19, OPW-01, OPW-02,           OPW-10, OPW-11, OPW-12, OPW-13, OPW-           15, OPW-18, OPW-19, AM773371 and		generated by primer OPW 05. Among 37	
OPM-09, OPM-11, OPM-13, OPM-14, OPM- 17, OPM-18, OPM-19, OPW-02, OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		primers only 20 primers (OPM-03, OPM-07,	
17, OPM-18, OPM-19, OPW-01, OPW-02,         OPW-10, OPW-11, OPW-12, OPW-13, OPW-         15, OPW-18, OPW-19, AM773371 and		OPM-09, OPM-11, OPM-13, OPM-14, OPM-	
OPW-10, OPW-11, OPW-12, OPW-13, OPW- 15, OPW-18, OPW-19, AM773371 and		17. OPM-18. OPM-19. OPW-01. OPW-02	
15, OPW-18, OPW-19, AM773371 and		OPW-10 OPW-11 OPW-12 OPW-13 OPW-	
		15. OPW-18. OPW-19 AM773371 and	
	<u> </u>		

AM750045) generated 100% polymorphic	
nroduets	
Constin diversity enclusio	
• Genetic diversity analysis	
<i>1. arjuna</i> accessions were grouped in three	
major clusters. In cluster-1, accessions 209, 123	
and 135 formed one sub-cluster. Out of them	
two are from Jharkhand and one from the	
neighboring State, Orissa. The other sub-cluster	
in the first cluster comprised of accessions 307,	
211 and 302, where two of them were from	
Maharashtra and one from Jharkhand. In the	
Cluster-II accession 512 (from Jharkhand) was	
placed as an isolate. One sub-cluster comprised	
of accessions 332 and 333, one from Andhra	
Pradesh and other from Chhattisgarh. The other	
sub-cluster comprised of three accessions, two	
(508, 430) from Jharkhand and one (504) from	
Andhra Pradesh. The third cluster had two sub-	
clusters and the first one comprised of 533 (UP)	
and 622 (Uttaranchal). Accession 701, 702 and	
703 (all from Maharastra) and 624 (from	
Uttarakhand) formed the second sub-cluster. The	
bootstrap values were significant except one i.e.	
40 for accessions 123 and 135. In comparison to	
the cluster analysis done for phenotypic and	
quantitative traits the number of clusters was	
less for RAPD data indicating that the	
genotypes are more influenced by environmental	
factors	
Three major aluster ware cheered for the	
intee major cluster were observed for the	
sixteen accessions of 1. tomentosa studied	
Cluster I comprised of seven accessions where	
accession 501 (Jharkhand) remained as isolate.	

	Accessions 309, 128 (both from Jharkhand), 216	
	(Maharashtra) and 229 (Chhattisgarh) comprised	
	of one sub cluster while other sub cluster had	
	accessions 310 (Chhattisgarh) and 408	
	(Jharkhand). The second major cluster showed	
	accessions 313 and 443 in sub-cluster and both	
	are from Chhattisgarh. Accessions 409	
	(Maharashtra) and 438 (Jharkhand) formed other	
	sub-cluster. Major cluster-III comprised of five	
	accessions, 522, 535 and 610 (all from	
	Jharkhand), 531 (Maharashtra) and 612 (Andhra	
	Pradesh).	