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Telah melaksanakan penelitian dengan judul sebagai berikut :

No	Judul Karya Ilmiah	Tahun Pelaksanaan
1	In the second state of Diach Linter Oil (Nigella section) as	renentian
1	Immunomodulatory Activity of Black Jinten Oli (Nigelia sativa) as	2020
	Macrophage Activator for Salmonella typimurium infected Rat	
2	Screening the Reproductive Tract of Dairy Cattle for Pathogenic	2019
	Micros	
3	Human Chorionic Gonadotropin (hCG) from Urine of Pregnant	
	Women to Manipulate in vivo Ovulation and Pregnancy of Madura	2019
	Cows	
4	Anti Early Embryonic Protein (EEP) for Pregnancy Test by	2010
	Microtiter Strip in Dairy Cows	2019
5	The Effect of Feeding High Level of Protein on Reproductive	2010
	Performance of Bali Starling.	2019
6	Antisperm Antibody in Repeat Breeder Friesian Holstein Cows at	
	KPSP Setia Kawan Nongkojajar, Tutur District, Pasuruan,	2019
	Indonesia.	
7	Diagnosis of Single and Twin Pregnancy, and Early Embryo	2010
	Mortality Through Progesterone Level Test on Local Does.	2019
8	Improvement of Pregnancy Rate in Bali Cows with the	
	Combination of Equine Chorionic Gonadotropine (eCG) from Local	2019
	Pregnant Mare with PGF2a.	
9	Progesterone Profile of Dairy Cows which Experienced the Failure	2010
	of Pregnancy to Artifical Insemination (AI).	2019
10	Effect of Heat Shock Protein (HSP) in Post Thaw Baluran Bull	2010
	Semen	2018
11	Potency of Mycotoxin Binders on MDA Level, Expressions of	
	Caspase 9 and Caspase 3 in The Uterus of Mice Exposed to	2017
	Zearalenone	















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12	Polymorphism of Growth Hormone Gene in The Artificial Insemination Result of Madura Cattle with Limousin Semen as a Reference for Canatia Selection	2018
12	Implementation of fate group the herizon of herizon to the second structure of	
15	of indo-pacific bottle nose dolphin (Tursiops aduncus) in bali	2020
	dolphin lodge	
14	Uji Sensitivitas Kebuntingan Sapi Perah Menggunakan Pregnancy	
	Specific Protein B (PSPB) Microtiter Strip dan Progesteron sebagai	2007
	Gold Standard	
15	Estimation of Equine Chorionic Gonadotropin (eCG) concentrate in	2014
	the Blood Sera of Pregnant Mare	2014
16	Efek Pemberian L-Arginin Terhadap Gambaran Histologi Jumlah	
	Spermatosit Primer pada Mencit (Mus musculus) Setelah Terpapar	2019
	Suhu Panas	
17	Anti Prolactine Overcomes Heat Stress on Laying Hen.	2008
18	Unnatural Forced Moulting in The Laying Hen as Cause of	2000
	Zoonosis from Salmonella Enteritidis	2009
19	Case Study: Dystocia on Beef Cattle in Kunir Regency of Lumajang	2017
	District, East Java, Indonesia in 2015 and 2016	2017
20	Teratogenic Effect of Congenital Toxoplasmosis in Chicken	2017
	Embryo	2017

Adapun penelitian tersebut layak dilakukan, meskipun belum ada *Ethical Clearence* karena menggunakan hewan coba yang minimal dan menghasilkan output yang sangat baik.

Demikian surat keterangan ini kami buat untuk dapat dipergunakan sebagai persyaratan pengususlan Jabatan Fungsional <u>Guru Besar</u>

Surabaya, 3 April 2023 、 Wakil Dekan III,

Prof. Dr. Mustofa Helmi Effendi, drh., DTAPH NIP 196201151988031002















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THE INDIAN VETERINARY JOURNAL

(Official organ of the Indian Veterinary Association)

Vol. 96

September 2019

CONTENTS

GENERAL ARTICLES :

Study on Morphological Traits in Indigenous <i>Jallikattu</i> Bulls of Tamil Nadu		
R. Priyadharsini, A. Gopinathan, S.M.K. Karthickeyan and P.N. Richard Jagatheesan		09
Epitopes Prediction According to Glycoprotein Encoding Gene of Rabies Virus Local Isolates as		
Vaccine Candidate against Circulating Rabies Virus in Indonesia		
JolaRahmahani, Suwarno and Fedik Abdul Rantam		14
Estimates of Genetic Parameters of Semen Traits in Crossbred Bulls		
Gopinathan, S.N. Sivaselvam, S.M.K. Karthickeyan and R. Venkataramanan		17
Commercial Cuts in Etawa Goat Fed with Water Spinach Pufa and Mineral Mixture Licks		
Kadek Rachmawati, Romziah Sidik, Vivi Virgianty, Koesnoto Supranianondo,		
Tri Bhawono Dadi and Herinda Pertiwi		20
Occurance of Ectoparasites in Mud Crab (Scylla serrata) and White Shrimp (Litopenaeus vannamei))	
Widyo Witular, Kismiyati and Putri Desi Wulansari		23
Antiviral Activity of Ethanolic Extract of Srikaya Seeds (Annona squamosa L.)		
Against Avian InfluenzaVirus		
Maya Nurwartanti Yunita, Adi Prijo Raharjo, Prima Ayu Wibawati and Bodhi Agustono	•••	26
Identification of Aeromonas hydrophila Outer Membrane Protein as a Potential Vaccine		
Candidate for Ulcer Disease		
M. Gandul Atik Yuliani, Anwar Ma'ruf, Ratna Damayanti and Diyantoro	•••	29
Diatom Naviculoid and Nitzschioid Composition in the Sulfur Mountains		
Watershed East Java, Indonesia		. .
Suciyono, A.A. Sabil, E. D. Masithah, D.D. Nindarwi, M.H. Azhar and M.F.Ulkhaq	•••	31
Anti Early Embryonic Protein (EEP) for Pregnancy Test by Microtiter Strip in Dairy Cows		
Abdul Samik and Erma Safitri	•••	37
Enhancement of Broiler Chicken Growth by Laserpuncture Treatment		
Lilian Soekwanto, Chairul Anwar and Rimayanti	•••	40
Protective Effect of Mycotoxin Binders on Ovarian Gestation Mice Exposed by Zearalenone		
Ragil Angga Prastiya, Abdul Samik, M. Thohawi Elziyad Purnama and Amung Logam Saputro	•••	44
Detection of Antibiotic Residue in Broiler Chicken Meat at Traditional		
Market in Surabaya City, Indonesia		
Shelly Wulandari, Diyantoro and OkySetyo Widodo	•••	47
The Potency of Hylocereus polyrhizus Peel Extract as Protector on Lead Acetate-Induced		
Testicular Toxicity in Mice		40
E. Wulandari, R. I'tishom R and S.A. Sudjarwo	•••	49
Probiotic Utilization in Megacolona Dog : A Case Report		
I ri BnawonoDadi, Ira Sari Yudaniayanti, WiwikiMisacoYuniarti, Nusdianto I riakoso,		F 4
BoediSeliawan, E. Djoko Poetranto, MilyayuSonetaSotyan and Herinda Pertiwi	•••	51
Honey as an Alternative to Stem Cells Therapy for Degenerated Rat Testis Due to Malnutrition		
Erma Satitri, Thomas V. Widiyatno, Willy Sandhika and R. Heru Prasetyo		53

CLINICAL AND FIELD ARTICLES :

Pathology of Selenite Induced Cataract in Rats		
S.Shiyamala, S.Ramesh, S.Hemalatha, C.Ramani, V.Ranganathan, K.Vijayarani,		
S.Ramesh and Ganne Venkata Sudhakar Rao		56
Successful Management of Recurrent Cervico Vaginal Prolapse in a Jersey Crossbred Cow by Caslick's Operation		
S. Rangasamy, T. Sarath, K. Logapriya, Aparna Gopinathan and J. Umamageswari		57
Dystocia Due to Hydrocephalus Fetus in a Queen Cat		50
S. Rangasamy, J.Umamageswari, Aparna Gopinatnan and D. Gopikrishnan	•••	59
Dystocia Due to Ringwomb in a Non Descript Doe S. Rangasamy, J.Umamageswari, Aparna Gopinathan and D. Gopikrishnan		61
Congenital Polycystic Kidney Disease in a Domestic Short Haired Cat- A Rare Case P.Ramesh, D.Sumathi, K.Jevaraja, C.Javanthi and M.G.Javathangaraj		63
Menaesonbanus in Dons – A Case Study		00
P.Ramesh, S.Kavitha, K.Jeyaraja, D.Chandrasekaran, S.Arun Prasad and M.G.Jayathangaraj		65
Therapeutic Management of Generalized Tetanus in an Adult Dog Due to Nail Bed Infection S. Sarayanan, M. Sasikala, K.M. Palanivel and G. Vijaya Kumar		67
Successful Management of Illogrative Pododermatitis in Rabbits with		0.
Sulfacetamide Dry Dusting Powder		
D.Divvalakshmi, N. Kumarvelu, Thanga, Thamil Vanan and P.Tensingh Gnanarai		69
Clinical Management of Oedematous Emphysema in HF Bull: A Case Report		
Vinod Haribhau Shende, Ajeet Singh, Anil Korade, S. Hanumantrao. Sontakke,		
Hemant Dashrath Kadam and Jayant Ramchandra Khadse		70
Ultrasonographic Evaluation of Gall Bladder Affections in Dogs - A Retrospective Study		
D.Sumathi, P.Ramesh, K.Jeyaraja, M.Ranjith Kumar and M.G.Jayathangaraj		72
Osteodystrophia Fibrosa in a Kathiyawari Colt - A Case Report		
R. Eazhisai, B. Gowri, A. Abinaya and S. Hamsa Yamini		74
Concurrent Clinical Coccidiosis and Subclinical Babesiosis in an Adult Cattle		
S. Saravanan, T. Mohanapriya, K.K. PonnuSwamy, P.A. Enbavelan,		
R.C. SundaraRajan and R. Ramprabhu		76
Diagnosis of Lymphoid Leucosis in an Organised Aseel Farm		
A.Varun, C. Yogesh, P.V.Sangeetha, S.Ezhilvalavan, M. Thangapandiyan, C.Pandian and A.V.Omprakash		78
Kaempferia galanga L. Inhibiting Effect on Vascular Endothelial Growth Factor (VEGF) and		
Cyclooxygenase-2 (Cox-2) Expression on Endothelium of Chorioallantoic Membrane		
Iwan Sahrial Hamid, Juni Ekowati and Muhammad Thohawi Elziyad Purnama		80
Pathology of Snake Envenomation in a Cat		
M.S.Sreelakshmi, Mammen, J. Abraham, N.D. Nair, B.Abeena and R.B.Vishnurahav		82
Black Quarter in a Crossbred Bull		_
M.S. Sreelakshmi, Ajith Jacob George, Mammen J. Abraham and N.Divakaran Nair		84
Author Index		87

Anti Early Embryonic Protein (EEP) for Pregnancy Test by Microtiter Strip in Dairy Cows

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(Received : March, 2019 78/19 Accepted : May, 2019)

Abstract

The purpose of this study was to identify and isolate of EEP from cotyledons of pregnant dairy cow for detecting early pregnancy by microtiter strip method in post-inseminated cows at 28 day, including confirmation by serum progesterone and utrasonography. The EEP with a molecular weight of 59.88 kDa could induce the onset of anti-EEP, which can be used for pregnancy test. The EEP microtiter strip test with blood serum progesterone and optimum USG levels was found sensitivity and specificity, as well as had positive predictive values and negative predictive values of 93.33%, 100%, 100% and 83.33%, respectively.

Keywords : early embryonic protein (EEP), Pregnancy test, Microtiter strip, Dairy cow

An early pregnancy diagnosis is a necessary step for identification to reduce the loss of production time as a result of infertility. This can be conducted in two ways: specifically by detecting specific substances contained in the parent's blood such as Early Embryonic Protein (EEP) (El-Amiri *et al.*, 2004) and by detecting non-specific substances in blood, urine or milk during pregnancy such as progesterone, estrone sulphate (Hafez, 2000).

Materials and Methods

The method of this research was used 6 stages : 1. Isolation EEP from cotyledon 60 days pregnant cattle and characterization by SDS-PAGE method, 2. Purification with Elution method (Hermadi *et al.*, 2018), 3. The making anti EEP in 6 male rabbits were injected subcutaneously with 150 µg of EEP isolates in *complete adjuvant Freunds*. The next booster was carried out 4th weeks after the initial injection with 100 µg of EEP isolates in *Freund's incomplete adjuvant*. 5 cc of blood was taken from each rabbit after 4th week, 4. The anti EEP was analyzed using indirect ELISA, 5. Anti-EEP was purificated using the Saturated Ammonium Sulphate (SAS) method, 6. The use of Anti-EEP for cow pregnancy diagnosis by microtiter strips method were designed based on the ELISA sandwich in post inseminated cows at day 28 including confirmation by serum progesterone by enzyme Immunoassay (EIA) and USG at 30th day after insemination.

Results and Discussion

After isolation and identification of EEP protein from cotyledon of pregnant dairy cows with immunohistochemistry, SDS-PAGE and western blotting, the molecular weight of EEP protein 59.88 kDa was obtained, as shown in Fig 1.

The results of the examination of EEP protein levels after artificial insemination can be seen in Table I. The results of the pregnancy examination using an USG can be seen in Fig 2.



Fig 1. Results of Western Blot EEP Protein in Pregnant Dairy Cattle Cotyledons

¹Corresponding author : Email : rma_fispro@yahoo.com

	Days after Artificial Insemination							
Cow	7		14		21		28	
	Abs	PL	Abs	PL	Abs	PL	Abs	PL
1	0.313	6260	0.470	9400	0.783	15660	0.563	11260
2	0.346	6920	0.519	10380	0.692	13840	0.554	11080
3	0.332	6640	0.532	10640	0.796	15920	0.598	11960
4	0.257	5140	0.463	9260	0.720	14400	0.514	10280
5	0.284	5680	0.568	11360	0.738	14760	0.483	9660
Average	0.306	6128	0.510	10208	0.746	14916	0.542	10848

Anti Early Embryonic Protein ...

Table I. EEP protein levels of blood in serum dairy cow after artificial insemination.

Abs: Absorbance; PL: Protein Level (µg / ml)



Fig 2. Pregnancy ultrasound results in 28-day pregnant dairy COW.

The results of the pregnancy diagnosis in dairy cows using the microtiter strip method, blood serum progesterone and ultrasound, were compared using the validity test (sensitivity and specificity) as shown in Table II.

The western blot results indicate that EEP molecules specifically bind to anti-EEP as primary antibodies and anti-rabbit IgG as secondary antibodies. Anti-EPF and anti-rabbit IgG could recognize EEP protein as a band with a molecular weight of 59.88 kDa. The protein band was in accordance with molecular weight of early embryonic protein (EEP), known as Pregnancy Associated Glycoprotein (PAG) in ovine placenta. 60-100 days of gestation found by El-Amiri et al. (loc cit) ranged from 55 to 66 kDa, whereas Garbavo et al. (1998) found the molecular weight of PAG in goat placenta was 55 kDa, 59 kDa and 62 kDa.

Austyn and Wood (1993) stated that to produce antibodies, repeat immunization can be carried out in rabbits. The lowest anti-EEP titre was obtained in week 5, which was increased after the booster. This was in line with the findings of Darnell et al. (1990) who stated that over a period of 2 weeks after injecting antigens, the concentration of IgG in serum began to rise, and quickly increased in the third week after the booster.

Pregnancy diagnosis with EEP microtiter strips were carried out at day 28 day after artificial insemination. To strengthen the success of the pregnancy diagnosis, blood serum progesterone levels were measured, and an

Table II. Pregnancy Diagnosis of Dairy Cows with EEP Strip Microtiter at 28 Post-IB Days and Progesterone Levels and USG at 28 Post-IB Days

Dragostarana		Results of Strip Microtiter				
Progesterone	036	Yellow (pregnant)	Colorless (Not pregnant)			
Pregnant		14	1			
Not Pregnant		0	5			
Total		14	6			

Sensitivity: 14:15 = 93.33%; Specificity: 5:5 = 100%

Positive predictive value: 14: 14 = 100%; Negative predictive value: 5: 6 = 83.33%

Abdul Samik and Erma Safitri

ultrasound was carried out. At day 28 after artificial insemination, the results of each of the 14 microtiter strips from 20th cows were positive for pregnancy. Xie et al. (1996) detected EPF in goats three weeks into the pregnancy. EPF concentration was reported to reach optimal before birth. Gonzales et al. (2000) found that the goat EPF concentration reached a maximum at 8 weeks gestation and began to decline over the period of 12-14 weeks, which remained the constant until delivery. Meanwhile, Vandaele et al. (2005) succeeded in detecting this substance at three to four weeks of the pregnancy, the concentration of EPF was directly proportional to the breed, fetus number and weight, while the age of the parent and sex of the fetus did not affect the EPF content. It was also reported that EPF concentrations began to decline dramatically after four weeks post partum.

The results of the progesterone blood serum testing on 28 dairy cows after artificial insemination concluded that 15 of the sample had progesterone levels > 1 ng / ml and 5 tails had a progesterone level <1 ng / ml. The progesterone levels of the pregnant dairy cattle obtained in this study was between 7.18 - 7.75 ng / ml.

Early embryonic protein during pregnancy plays a role in maintaining the *corpus luteum* by stimulating the production of prostaglandin E2 (PGE2). Prostaglandin E2 will increase cAMP synthesis so that the progesterone production in the corpus *luteum* will also rise during pregnancy (Karen *et al.*, 2001).

The results of the blood serum progesterone test, ultrasound and EEP Microtiter strips revealed sensitivity, specificity, positive predictive value and negative predictive value at rates of 93.33%, 100%, 100% and 83.33% respectively, for pregnancy.

Summary

The conclusion, Pregnancy testing with a microtiter strip, 28 days after artificial insemi-

nation is considered both valid and accurate because during that period of time, the placenta is formed, hence the EEP protein can be released into the maternal peripheral blood. It is therefore detected with an anti-EEP in the strip microtiter.

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THE INDIAN VETERINARY JOURNAL

Vol. 96

September 2019

No. 09

AUTHOR INDEX

Abdul Samik	37,	44
Abeena, B.		82
Abinaya, A.		74
Adi Prijo Raharjo,		26
Ajeet Singh,		70
Ajith Jacob George,		84
Amung Logam Saputro		44
Anil Korade,		70
Anwar Ma'ruf,		29
Aparna Gopinathan 57,	59,	61
Arun Prasad, S.		65
Azhar, M.H.		31
Bodhi Agustono,		26
BoediSetiawan,		51
Chairul Anwar		40
Chandrasekaran, D.		65
Divakaran Nair, N.		84
Divyalakshmi, D.		69
Diyantoro,	29,	47
Djoko Poetranto, E.		51
Eazhisai, R.		74
Enbavelan, P.A.		76
Erma Safitri	37,	53
Ezhilvalavan, S.		78
Fedik Abdul Rantam		14
Gandul Atik Yuliani, M.		29
Ganne Venkata Sudhakar R	ao	56
Gopikrishnan, D.	59,	61
Gopinathan, A.	09,	17
Gowri, B.		74
Hamsa Yamini, S.		74
Hanumantrao. Sontakke, S.		70
Hemalatha, S.		56
Hemant Dashrath Kadam		70
Herinda Pertiwi,	20,	51
Heru Prasetyo, R.		53
l'tishom R,		49
Ira Sari Yudaniayanti,		51
Iwan Sahrial Hamid,		80

Javant Ramchandra Khadse	70
Javanthi, C.	63
Jayathangaraj, M.G. 63, 65	, 72
Jeyaraja, K. 63, 65	, 72
JolaRahmahani	14
Juni Ekowati,	80
Kadek Rachmawati,	20
Karthickeyan, S.M.K. 09	, 17
Kavitha, S.	65
Kismiyati,	23
Koesnoto Supranianondo,	20
Kumarvelu, N.	69
Lilian Soekwanto,	40
Logapriya, K.	57
Mammen J. Abraham, 82	, 84
Masithah, E.D.	31
Maya Nurwartanti Yunita,	26
MiyayuSonetaSofyan,	51
Mohanapriya, T.	76
Muhammad Thohawi Elziyad	
Purnama,	80
Nair, N.D.	82
Nindarwi, D.D.	31
NusdiantoTriakoso,	51
OkySetyo Widodo	47
Omprakash, A.V.	78
Palanivel, K.M.	67
Pandian, C.	78
PonnuSwamy, K.K.	76
Prima Ayu Wibawati,	26
Priyadharsini, R.	09
Putri Desi Wulansari,	23
Ragil Angga Prastiya,	44
Ramani, C.	56
Ramesh, P. 63, 65	, 72
Ramesh, S.	56
Ramprabhu, R.	76
Ranganathan, V.	56
Rangasamy, S. 57, 59	, 61

Ranjith Kumar, M.		72
Ratna Damayanti,		29
Richard Jagatheesan, P.N.		09
Rimayanti		40
Romziah Sidik,		20
Sabil, A.A.		31
Sangeetha, P.V.		78
Sarath, T.		57
Saravanan, S.	67,	76
Sasikala, M.		67
Shelly Wulandari,		47
Shiyamala, S.		56
Sivaselvam, S.N.		17
Sreelakshmi, M.S.	82,	84
Suciyono,		31
Sudjarwo, S.A.		49
Sumathi, D.	63,	72
SundaraRajan, R.C.		76
Suwarno		14
Tensingh Gnanaraj, P.		69
Thanga. Thamil Vanan,		69
Thangapandiyan, M.		78
Thohawi Elziyad Purnama, M	Л.	44
Thomas V. Widiyatno,		53
Tri Bhawono Dadi	20,	51
Ulkhaq, M.F.		31
Umamageswari, J. 57,	59,	61
Varun, A.		78
Venkataramanan, R.		17
Vijaya Kumar, G.		67
Vijayarani, K.		56
Vinod Haribhau Shende,		70
Vishnurahav, R.B.		82
Vivi Virgianty,		20
Widyo Witular,		23
Willy Sandhika,		53
WiwikMisacoYuniarti,		51
Wulandari, E.		49
Yogesh, C.		78

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