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PION OR OT ETYAL DECAMINACE

The 4th International Anatomical Sciences and Cell Biology Conference

4th - 6th December, 2016 (Sun - Tue) | Hong Kong

CONFERENCE VENUE

Cheung Kung Hai Conference Centre Li Ka Shing Faculty of Medicine The University of Hong Kong

> This conference is part of the 130th Anniversary Frontiers Conference Series in celebration of 130 Years of Medicine in Hong Kong.



Schedule-at-a-Glance

advenin

Super-resolution & Intra-vital Imaging Workshop December 4, 2016 (Sunday)

08:30 - 17:00	Registration	
Time	Programme	Venue
09:30 - 11:00	L1-L3	Lecture Theatre 3
11:00 - 11:30	Coffee Break	Exhibition Area
11:30 - 12:30	L4 – L5	Lecture Theatre 3
13:00 - 14:00	Lunch	Exhibition Area
14:00 - 18:00	Demonstration Session 1 - 4	FCF

Conference – Day 1 December 5, 2016 (Monday)

Time	Programme			
08:30 - 17:30	Registration			
	Lecture Theatre	4		
09:00 - 09:20	Opening Ceremony Guest of Honor: Professor LEUNG, Suet-yi Associate Dean (Research), LKS Faculty of Medicine The University of Hong Kong, Hong Kong			
99; 20 – 10 : 50	Lecture Theatre 4 Plenary Lecture 1			
10:50 - 11:15	Coffee Break / Poster Session			
	Lecture Theatre 3	Lecture Theatre 4		
11:15 – 12:45	SYM 1.1 Advances in Neuroscience	SYM 1.2 Innovations in Teaching Histology		
12:45 - 14:00	Lunch Break / Poster Session			
	Lecture Theatre 3	Lecture Theatre 4		
14:00 - 15:30	SYM 2.1 Advances in Developmental Studies	SYM 2.2 Good Practices in Teaching Gross Anatomy		
15:30 - 16:00	Coffee Break / Poster Session			
16:00 - 17:30	Lecture Theatre 4 Plenary Lecture 2			
18:30 - 21:00	Banquet (Full registration participants and in Golden Lilies Banquet, Cyberport, 1			

Conference – Day 2 December 6, 2016 (Tuesday)

Time	Pro	gramme
08:30 - 14:00	Registration	•
09:00 - 10:30	Lecture Theatre 4 Plenary Lecture 3	
10:30 - 11:00	Coffee Break / Poster Session	
	Lecture Theatre 3	Lecture Theatre 4
11:00 - 12:30	SYM 3.1 Advances in Cancer Research	SYM 3.2 Body Donation
12:30 - 13:45	Lunch Break / Poster Session	
	Lecture Theatre 3	Lecture Theatre 4
13:45 – 15:15	SYM 4.1 Advances in Aging Studies	SYM 4.2 Innovations in Teaching Gross Anatomy
15:15 - 15:45	Coffee Break / Poster Session	
15:45 – 17:15	Lecture Theatre 4 Plenary Lecture 4	
17:15 - 17:30	Closing Ceremony / Poster Awards	

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Poster Abstract 45 Developmental Biology

Correlation between Interleukin-17 Serum and Goiter Gradation of Graves' disease patients

Dwitya Elvira, Eryati Darwin Internal Medicine Department of Medical Faculty of Andalas University, Indonesia

s' disease (GD) is a thyroid autoimmune disease that systemically affect human metabolism. Graves' disease exercized as hyperthyroidism, diffuse goiter, opthalmopathy and/ dermopathy, increasing thyroid hormone decrease of thyroid stimulating hormone (TSH) and increasing of autoantibody thyroid (TRAb). of GD remains elusive but imbalance of T-helper cells, regulatory T cells and cytokines were accused to this disease. The identification of new subpopulation of T helper (Th-17) produce IL-17 (Interleukin-17), crucial in development of autoimmune disease, particularly Graves' disease. Gradation of goiter were anto three classifications: grade I, grade II and grade III. Aim of this study is to correlate between serum of cytokine with goiter gradation of Graves' disease patients. Thirty patients diagnosed as Graves' disease on clinical and laboratory measurement were included as object of this study. Age, gender, goiter gradation, function status was measured and noted as baseline characteristic of this study. Goiter gradation classified classifications: grade I, grade II and grade III. Serum of IL-17 were measured using ELISA method. As a of this study, we found increasing of IL-17 cytokine serum compared to control, with mean values 11,961,73 m patient group, and 7,477,07 pg/mL in control group. Goiter gradation found mostly in grade II (50%) by grade I and grade III. There is no correlation between IL-17 serum with goiter gradation. From this we can conclude that there is no correlation between cytokine IL-17 serum with goiter gradation in Graves' patients but we found increasing of interleukin-17 serum in Graves' disease patients that suggested that cells can play a central role in pathogenesis of Graves' disease.

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Effect of Fixed Orthodontic Appliance on the Levels of Matrix Metalloproteinase-8 in Gingival Crevicular Fluid

Eryati Darwin* and Aida Fitriana**

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orthodontic therapy is the preferred therapeutic modality for treatment of malocclusions that affect social appearance. Nevertheless Its may cause subgingival microbial composition, promotion of development and soft tissue injuries. This mechanical injury can lead to development of inflammatory in periodontal tissues, which promote tooth movement and biological processes, resulting in bone toon. Matrix metalloproteinases (MMP) are thought to be responsible for the turnover and degradation of the seculular matrix that act on pro-inflammatory cytokines. Matrix metalloproteinase-8 (MMP-8) is a member of family of enzymes is seen in almost every human tissue in which inflammation is present.

healthy patients who treated with a straight wire technique using brackets on the maxillary and the healthy patients who treated with a straight wire technique using brackets on the maxillary and the healthy patients were divided into two group; one group with less than six months treated, and the other group more than six months treated. Salivary samples were taken from gingival crevicular fluid and its MMP-8 were determine using ELISA method. This study was approved by Research Ethic Committee.

study shows that the level of MMP-8 in group using more than six months fixed orthodontic appliance were than in group using less than six months with statistically significant differences (p<0,05).

our study, we conclude that orthodontic appliance may induced inflammatory process on the gingival tissue

words: fixed orthodontic, inflammation, MMP-8

Targeting ANXA3 in combination with sorafenib for the treatment of hepatocellular carcinoma

Man TONG, Steve LUK, Jin DING, Terence LEE, Stephanie MA School of Biomedical Sciences, LKS Faculty of Medicine, The University of Hong Kong

Sorafenib is the only FDA-approved tyrosine kinase inhibitor for targeted therapy in advanced HCC. Neverther its efficacy is limited with only a modest improvement in patient outcome, likely due to acquired resistance. depth understanding of the molecular mechanism of sorafenib resistance is warranted for the development of sorafenib resistance is so that the development of the d treatment strategies. Recent studies by us and others have characterized liver tumor-initiating cells (T-ICs) possible source of resistant and recurrent tumors and a plausible target for HCC treatment. Our group previously identified CD133 to be a functional marker of liver T-ICs and found annexin a3 (ANXA3) to recommend cancer and stem cell-like properties in this subset of cells. Interestingly, our recent observations also found liver T-ICs to be more resistant to sorafenib. Sorafenib resistant clones, established in HepG2 and Huh7 continuous exposure to increasing concentrations of sorafenib, displayed enhanced abilities to migrate. self-renew, and initiate tumor formation in immunodeficient mice, as well as higher expression of associated genes. These two sorafenib resistant cell lines and two other sorafenib resistant HCC patientxenografts established in a similar manner were also found to be enriched for CD133 and ANXA3 expenses Sorafenib resistant clones with ANXA3 stably suppressed were re-sensitized to sorafenib treatment and sensitized treatment and sensit diminished ability to migrate, invade, self-renew and initiate tumor growth in vivo, further substantiating of ANXA3 in mediating sorafenib resistance in HCC. Mechanistically, an activated PKC/ERK/FRA2 axis was found to be responsible for driving this phenomenon. Clinically, ANXA3 expression was also found have prognostic value as a higher ANXA3 expression in HCC patients who have received sorafenib treatments correlated with poor overall survival. The combinatorial use of a homemade ANXA3 neutralizing antibases sorafenib on HCC patient derived xenografts is now being investigated as a potential new treatment reg combating sorafenib resistance in HCC.

Developmental E

The Correlation of Prolactin Level and Oxytocin with Duration of Amenorrhea Lactation in Exclusive Breastfeeding Women

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¹Department of Biology

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Faculty of Medicine, Andalas University, Indonesia

Prolactin and oxytocin were hormones that play important roles in lactation process. When mothers expenses breastfeeding, it presses ovulation causing no occurrence of menstruation, called amenorrhea lactation. At months breastfeeding, it would make the mother had higher risk for menstruation and pregnancy. The arm research is determining the correlation between prolactin and oxytocin level with duration of amenorrheading exclusively breastfeeding mothers.

The design of the research was cross-sectional, observing 48 exclusively breastfeeding mothers in Padang Belimbing, September 2015 - June 2016. Samples were selected based on cluster random samples examination of prolactin and oxytocin level was conducted at Biomedical Lab Faculty of Medicine United human prolactin ELISA Kit and Human oxytocin Elisa Kit. Shapiro-Wilk was applied for normality test data, and Spearman's correlation was applied for analyzing prolactin and oxytocin level with daragements.

The result is that there is a weak positive correlation and significance between prolactin level and durant amenorrhea lactation (r=0.331; p=0.022); and there is a very weak negative correlation and insignificance oxytocin level with the duration of amenorrhea lactation (r=-0.085; p=0.565).

The higher prolactin level, the longer amenorrhea lactation duration and the higher oxytocin level, the amenorrhea lactation period.

Keywords: Prolactin, Oxytocin, Duration of Amenorrhea Lactation, Exclusive Breastfeeding



Effect of Fixed Orthodontic Appliance on the Level of Matrix Metalloproteinase-8 in GingivalCrevicular Fluid

Eryati Darwin¹ and Aida Fitriana²
¹Faculty of Medicine Andalas University, ²Faculty of Dentistry Andalas University, Padang-Indonesia

Background

Fixed orthodontic therapy is the preferred therapeutic modality for treatment of malocclusions that affect mastication and facial appearance. Nevertheless its may cause subgingival microbial composition, promotion of plaque development and soft tissue injuries. This mechanical injury can lead to development of inflammatory reactions in periodontal tissues, which promote tooth movement and biological processes, resulting in bone resorption. Matrix metalloproteinases (MMP) are thought to be responsible for the turnover and degradation of the extracellular matrix that act on pro-inflammatory cytokines. Matrix metalloproteinase-8 (MMP-8) is a member of MMP family of enzymes is seen in almost every human tissue in which inflammation is present. To determine the effect of fixed orthodontic appliance, we measured the level of matrix metalloproteinase in gingival crevicular fluid



Material and Methods

- 10 subjects of healthy patients treated with a straight wire technique using brackets on the maxillary and the mandibular arches for less than 6 months and more than 6 months
- 10 subjects using brackets on the maxillary and the mandibular arches as accessories for less than 6 months and more than 6 months
- Salivary samples were taken from gingival crevicular fluid using paper point no.15
- MMP-8 level were measured by using ELISA method.
- Statistical analysis: Kruskall Wallis and Post Hoc Man Whitney

Results

- Kruskall Wallis analysis for the level of MMP 8 on gingival crevicular fluid of subjects were using fixed orthodontic appliance for treatment and accessories purpose for less than 6 months and more than 6 months sows in table
- The level of MMP-8 in group using more than six months fixed orthodontic appliance were higher than in group using less than six months with statistically significant differences (diagram).

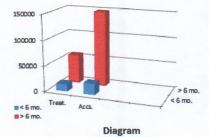






Table: The level of MMP 8 on gingival crevicular fluid of subjects were using fixed orthodontic appliance for treatment and accessories purpose for less than 6 months and more than 6 months

Fixed Orth. Appliance	MMP 8 level (pg/dl)	p
	Median	
Treatment <6 month	14237,56	0,148
Treatment > 6 month	56830,94	
Accessories < 6 month	21091,67	
Accessories < 6 month	147892,66	



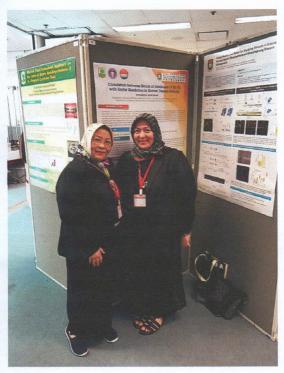
Conclusion

From our study, we conclude that long period of fixed orthodontic appliance may induced inflammatory process on the gingival tissueespecially for the long period fixed orthodontic appliance for accessories purpose

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Certificate of Attendance

This is to certify that

DARWIN Eryati

has attended the

4th International Anatomical Sciences and Cell Biology Conference (IASCBC 2016)

held in The University of Hong Kong on 4th-6th December, 2016.

Thur Sie Der

George Tsao Sai Wah Chairman of Organising Committee IASCBC 2016



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