

## *Editorial*

### **American Journal of Experimental and Clinical Research**

It is my great pleasure to announce the launch of the American Journal of Experimental and Clinical Research [ISSN 2330-9237 (Print) and ISSN 2330-9245 (Online) <http://www.ajecr.org>]. We have worked tremendously in getting this journal ready to launch and are confident that the journal will rapidly progress during a short period of time to reach a highly competitive level.

We welcome you to this new journal! The journal was founded by a group of scientists who are devoted to the advancement of experimental and clinical medical and dental research and their clinical applications.

During the past decades, considerable progress has been made in basic and clinical medical researches with the ultimate aim of understanding and unraveling the mechanism(s) underlying the development of human diseases. These achievements have provided challenges in eliminating or alleviating the effects of human illnesses and thus helping to prolong the life spans of human being. It was only several decades ago that people frequently died from general diseases which nowadays are considered highly treatable. Such diseases as influenza, tuberculosis or diarrhea no longer pose a significant mortality threat in contemporary society. Thanks to the discovery of powerful vaccines and also increased public health education level to prevent contagious or communicable diseases. Although other diseases such as heart diseases and cancers still account for primary cause of death, considerable improvements in the early diagnosis and treatment of such disorders have enabled increased survival and prolonged life of the patients.

With the invention of new technologies, medications and procedures, the effectiveness of the treatment of previously lethal diseases has dramatically increased. The availability of ultrasound, computerized tomography, magnetic resonance imaging, positron emission tomography (PET) and so on that are less invasive techniques has also increased the effectiveness of early diagnosis and thus more effective therapy. All of these achievements have been brought about through extensive experimental and clinical researches.

The impacts of experimental and clinical research on human life expectancy are undeniable and will be more pronounced with rapid advances in biomedical research. Currently, cellular and molecular research scientists are rigorously engaged in the development of methodologies and pharmaceuticals that will eradicate human diseases. The American Journal of Experimental and Clinical Research provide a platform for these studies to be shared with the research community at large.

The American Journal of Experimental and Clinical Research is an open-access, peer-reviewed online and print journal that encompasses all aspects of experimental and clinical research in the field of medicine and dentistry. The development of open-access online journals has created new opportunities and new challenges for publishers, authors, and readers.

The American Journal of Experimental and Clinical Research aims to publish articles contributing to the development of basic and clinical medical and dental science researches and also encourages submissions considering the new methodologies to facilitate experimental approaches. The journal is at present a quarterly published journal that is dedicated to be a high quality platform to facilitate rapid publication and circulation of novel discoveries in all aspects of basic and clinical medical and dental researches. The American Journal of Experimental and Clinical Research welcomes original and review articles on both clinical investigation and basic medical research. There are also categories such as case reports, short communications, methods/techniques, letter to the editor/correspondences and editorials.

We believe that the success and reputation of the American Journal of Experimental and Clinical Research depend on the quality of the articles published and therefore encourage you to submit your important research work to this journal. In addition, we believe that providing a fair and rapid peer-review and manuscript process to publication is of utmost importance to our authors. We also continuously improve our journal standing by publishing high standard articles.

We would like to take this opportunity to thank our editorial board members, authors and editorial staff for their support and active participation in making this journal possible. In addition, we would like to invite all of our colleagues to consider the American Journal of Experimental and Clinical Research as a future venue for their research works.

M. Ghazizadeh, MD, PhD  
Editor-in-Chief  
American Journal of Experimental and Clinical Research  
<http://www.ajecr.org>

## Original Article

# Variations in adrenal hormones in law enforcement servicemen during a mission to local armed conflict

Roman Koubassov\*, Yury Barachevsky

*Department of Medicine Catastrophe, Northern State Medical University, Archangelsk, Russia*

**Abstract.** In a previous study, we reported changes in the adrenocorticotropic hormone (ACTH) and cortisol secretion in blood samples from law enforcement personnel during the mission to local armed conflict region. In the present study, we demonstrate those changes collectively with additional data on changes in the adrenaline and noradrenaline in the urine samples of the same individuals. The study was conducted on 48 male officers who were deployed to an army conflict territory for a duration of 4 months. At the onset of the mission, there was a modest increase in all hormones corresponding to the general adaptation syndrome theory. As the mission started, significant increases were observed in the mean levels of the hormonal parameters in both serum and urine at different time points as compared to those before the mission. At first week of deployment, a sharp increase in the secretory activity of medulla and cortical adrenal gland was found and at the termination of the mission a dysfunction of hypophysis-adrenal gland regulation system was identified. These findings might lead to disturbances in interhormonal relationships and cause decreased stress tolerance in the relevant individuals.

**Keywords:** Law enforcement officer, adrenaline, noradrenaline, adrenocorticotropic hormone, cortisol, emergency condition

### Introduction

The principal responsibility of any government is to secure safety of its citizens. Accordingly, every type of life activity has to be protected from potential conflict hazards.

The world community has entered into a new era in the 21st century, creating intensification of different political, ideological, religion, or economical conflicts and crises. Some factors with much impact on social transformation includes technological progresses and global environment changes [1-3].

In order to maintain law enforcement in different territories and secure the safety of citizens, special police squads are needed. The professional task of law enforcement officers occurs in extreme conditions and often in emergency situations. The service duties are in the range of medium security with hardware assistance, special equipment and different weapons [4, 5]. In addition, besides professional detrimental factors affecting an armed personnel who has been trans-located from another region, other factors such as specific climatologic and geographical environments of a combat territory are pivotal [6, 7]. These factors based on their severity and duration may create considerable health problems ranging from functional disorders to pathological conditions with permanent impairments [8, 9]. In such individuals, various functional changes occur in order to provide adaptation to those conditions. In fact, the endocrine system plays a

major role in forming a compensatory regulatory mechanism to counter extreme impacts. In such response, activation of sympathoadrenal system plays a pivotal role [10, 11].

In a previous study, we reported changes in the adrenocorticotropic hormone (ACTH) and cortisol secretion in blood samples from law enforcement officers during the mission to local armed conflict area [12]. Subsequently, the aim of the present study was a comprehensive assessment of changes in ACTH, cortisol, adrenaline, and noradrenaline, secretions in both blood and urine samples from the same group of servicemen during a mission to the local armed conflict territory.

### Materials and Methods

We studied 48 male officers (mean age: 28.28±0.51) from the Ministry of Home Affairs who were law enforcement servicemen. All subjects had a mission to an army conflict territory (North Caucasus) for the purpose of maintaining law enforcement. The duration of their mission was 4 months. In all cases, we measured the blood serum ACTH by radioimmunoassay (Cis bio International, Cedex, France) and cortisol by enzyme immunoassay (Monobind Inc, California, USA). In addition, the levels of adrenaline and noradrenaline in urine samples of all cases were measured using gass chromatography method with mass spectrometer detector. Statistical analysis of the data was performed using the SPSS 15.0 software. The mean and

\* Corresponding author: Roman V Koubassov, PhD  
 (roman2001@gmail.com).

standard deviation (SD) was calculated for each measurement. To assess the universal distribution, Shapiro-Wilk normality test was applied. Comparative analysis of means was performed by Wilcoxon rank test. A P value less than 0.05 was considered as significant.

## Results

Overall the results of our study showed that in all cases, the levels of the hormones examined were in the normal physiological range before the mission. However, significant increases were found in the mean levels of the hormonal parameters at different time points during the mission as compared to those before the mission.

The mean serum ACTH level was increased more than two-fold in the individuals at 14 days after mission as compared to before mission (Table 1). At 1 month after mission, the ACTH level was increased four-fold ( $p < 0.001$ ). At 2 months after mission, we found a subsequent ACTH rise but it was less significant when compared with the measurement at 1 month ( $p = 0.005$ ). At the time of termination of the mission, the mean ACTH level was decreased but it still remained higher than the level before mission ( $p < 0.001$ ).

The serum cortisol level changed during the combat mission with a different pattern from ACTH dynamics (Table 1). Particularly, at 14 days after the mission, the mean cortisol level increased as compared with before mission ( $p = 0.002$ ). However, it was decreased at 1 month ( $p = 0.006$ ) and 2 months ( $p < 0.001$ ) as compared with 2 weeks after the mission. The mean cortisol level at 1 and 2 months showed no statistical difference as compared to before mission ( $p > 0.05$ ). At the end of the mission (after 4 months), the cortisol level increased to a maximum level. The level was similar to that of the two weeks measurement but better than before the mission and those at 1 and 2 months ( $p = 0.05$  and  $p = 0.003$  respectively).

The dynamics of urinary catecholamine levels in law enforcement servicemen during combat mission had analogous features with serum ACTH and cortisol changes but with less statistical significance (Table 2). Urinary excretion of adrenaline in 2 weeks combatants was increased twice as compared with before the mission ( $p < 0.001$ ). At 1 month, this parameter remained similar to that at 2 weeks ( $p = 0.65$ ). At 2 months, urinary adrenaline concentration reached the highest level as compared with 2 weeks and 1 month measurements ( $p = 0.002$  and  $p = 0.02$  respectively). When the combat mission was over, the mean adrenaline level decreased, but it was higher than that of before the mission ( $p = 0.04$ ).

The analyses of urinary noradrenaline excretion showed an increase at 2 weeks after the mission (Table 2). At 1 month after the mission, unlike adrenaline, the noradrenaline level increased as compared to that at 2 weeks ( $p = 0.05$ ) and continued at a similar level at 2 months. When the combat mission was over, noradrenaline concentration decreased, but it was higher than that of before the mission ( $p = 0.003$ ).

## Discussion

Suprarenal hormones play a leading role in the formation

TABLE 1  
SERUM ADRENOCORTICOTROPIC HORMONE (ACTH)  
AND CORTISOL CHANGES AT DIFFERENT DURATIONS  
OF MISSION

Duration	ACTH (pg/ml)	Cortisol (nmole/l)
Before mission <sup>a</sup>	20.07±3.20	404.81±124.54
2 weeks <sup>b</sup>	55.80±15.68	489.25±112.46
1 month <sup>c</sup>	86.43±17.56	426.80±102.54
2 months <sup>d</sup>	96.60±17.18	407.82±101.66
4 months <sup>e</sup>	55.37±10.70	471.16±117.78

Values are mean±SD. Statistical significances: ACTH (a vs. b,c,d,e, and b vs. c,d, and c vs. e and d vs. e,  $p < 0.001$ ; b vs. e,  $p < 0.81$ ; c vs. d,  $p < 0.005$ ). Cortisol (a vs b,  $p < 0.002$ ; a vs. c,  $p < 0.35$ ; a vs. d,  $p < 0.89$ ; a vs. e,  $p < 0.003$ ; b vs. c,  $p < 0.006$ ; b vs. d,  $p < 0.001$ ; b vs. e,  $p < 0.44$ ; c vs. d,  $p < 0.37$ ; c vs e,  $p < 0.05$ ; d vs e,  $p < 0.006$ ).

TABLE 2  
URINARY ADRENALINE AND NORADRENALINE  
CHANGES AT DIFFERENT DURATIONS OF MISSION

Duration	Adrenaline (nmole/day)	Noradrenaline (nmole/day)
Before mission <sup>a</sup>	83.01±18.63	160.35±38.85
2 weeks <sup>b</sup>	117.12±55.68	229.30±102.34
1 month <sup>c</sup>	122.80±67.56	275.68±121.58
2 months <sup>d</sup>	161.55±77.18	268.32±111.67
4 months <sup>e</sup>	91.32±57.62	193.61±66.05

Values are mean±SD. Statistical significances: Adrenaline (a vs. b,c,d, and d vs. e,  $p < 0.001$ ; a vs. e and a,c vs. e,  $p < 0.04$ ; b vs. c,  $p < 0.65$ ; b,c vs. d,  $p < 0.002$ ; b vs. e,  $p < 0.03$ ). Noradrenaline (a vs. b,c,d, and c,d vs. e,  $p < 0.001$ ; a vs. e,  $p < 0.03$ ; b vs. c,e,  $p < 0.05$ ; b vs. d,  $p < 0.07$ ; c vs. d,  $p < 0.76$ ).

of adaptation response to environmental factors in humans [13]. Abnormality of adrenal regulatory function occurs during excessive and long-term exposure to harmful environmental agents and results in a decline in human performance and physical and mental suffering (distress). The principal manifestations of these abnormalities are hormonal hypersecretion, target cell resistance, and failure of feedback regulation mechanism [14, 15].

Various investigations have shown that combatants with first time experience of deployment to a mission have allostasis laboratory markers of increased ACTH and cortisol levels after the mission. The level reaches to a maximum after 2 weeks. At the end of combat mission, these parameters decrease but remain higher than those at the beginning of the mission. This is indicative of a homeostasis imbalance retention [16]. A number of studies demonstrate that persistence of high levels of catecholamine, ACTH and cortisol in combatants for more than 6 months predetermine a triggering pathogenic mechanism of posttraumatic stress disorder [17-19]

In our study, the serum ACTH level at 2 months after the combat mission was constantly increased. At the time of termination of the mission (after 4 month), it decreased, but still exceeded twice when compared with that of before the mission. With regard to the serum cortisol changes, in spite of a sharp increase after the first two weeks, a decrease of this hormone was observed, although ACTH

was increased. Therefore, hormonal disbalance could be the first sign of disturbances in hypophysis-adrenal gland regulation system. At the end of mission, a dramatic cortisol increase was found, comparable with the first 2 weeks. In fact, this may predestine a derangement of adaptation process.

Catecholamine (adrenalitermine or noradrenaline) generated in adrenal medulla produces a short-term adaptation process. It has a catabolic effect and involves in almost all metabolisms. Catecholamine secretion is increased in stress conditions and extreme situations. The target cells and action mechanisms for adrenaline and noradrenaline are different though of common physiological effect. Thus, adrenaline (named as "fear hormone") improves individual's weakness resistance in initial stress time at high speed. The noradrenaline (named as "fury hormone") act after adrenaline. Its blood secretion corresponds with aggression and promote muscle strength. Noradrenaline is potent as adrenaline effect [20, 21].

In our study, we found that at first two weeks after the combat mission, the urinary adrenaline and noradrenaline levels were increased in the military servicemen. This observation could be attributed to the natural adaptation process to environmental changes. Furthermore, at two months, the adrenaline level was increased but when the mission was over it was decreased. However, the adrenaline level remained higher than that before the combat mission. This observation could be due to a retardation of adaptation process and probably first sign of distress. Concerning the noradrenaline changes, there was a significant increase and prolonged duration of this hormone. From the physiologic point of view, this appears to be a normal individual's response required for elongation of survival probability in stress situation. Similar results have been obtained in other investigations. Taken together, in the military servicemen during the armed mission a specific feature of catecholamine dynamics was observed which was attributable to adaptation process but in some cases with signs of distress. These hormonal imbalances could remain up to 6 month or more [22].

In conclusion, in combatants particularly special police squads that have acquired professional skills in extreme conditions and during the mission to local armed conflict territory, the secretory function of adrenal gland undergoes changes corresponding to the principles of general adaptation syndrome theory. At first week after the mission to military zone, a sharp increase is seen in the secretory activity of medulla and cortex of adrenal gland. At the termination of combat mission, signs of dysfunction in hypophysis-adrenal gland regulation system appear that may lead to disturbances in interhormonal relationships and thus weakening of stress tolerance.

The disturbance of endocrine regulation requires establishment of special measures to reduce it. The ultimate aim of these precautions is to provide increased resistance and capability to deal with extreme conditions in emergency case and prevention of mortality risk. These measures should include a long-range and clear-cut planning of combat missions, early diagnostics including

laboratory tests for armed personnels to be deployed to harmful locations, special training (physical, psychological, preventive, etc.) for armed personnels who are going to work in extreme conditions and emergency situations, and special medical rehabilitation measures that provide rapid restoration of the individual's health after the termination of combat mission. .

### Conflict of Interest

The authors declare no conflicts of interest.

### References

- Iverson T, Perrings C. Precaution and proportionality in the management of global environmental change. *Glob Environ Chang* 22:161-177, 2012.
- Kovats RS, Butler CD Global health and environmental change: linking research and policy. *Curr Opin Environ Sustain* 4:44-50, 2012.
- Schulze-Makuch D, Irwin L.N, Fairén A.G. Drastic environmental change and its effects on a planetary biosphere. *Icarus* 225:775-780, 2013.
- Brisebois R, Hennecke P, Kao R, McAlister V, Po J, Stiegelmar R, Tien H. Canadian Forces Health Services Research Consortium. The Role 3 Multinational Medical Unit at Kandahar Airfield 2005-2010. *Can J Surg* 54:124-129, 2011.
- Koubassov RV, Barachevsky YE, Lupachev VV. Problems of professional safety of local armed conflict servicemen. *Medico-Biological and Socio-Psychological Problems of Safety in Emergency Situations*. 1:39-46, 2014. (in Russian).
- Shellman SM, Hatfield C, Mills M.J. Disaggregating actors in international conflict. *J Peace Res* 47:83-90, 2010.
- Herrell RK, Bliese PA, Hoge CW. Effect of combat intensity, depression, alcohol misuse, and family history of depression and alcohol misuse on PTSD in a sample of post-deployment US Soldiers. *Compr Psychiatry* 54:e4-e5, 2013.
- Artiss K. The combat soldier. *Mil Med* 165:33-40, 2000.
- Dobson M. Combat Stress Reaction. In *Encyclopedia of Stress* (Fink G. ed). USA: Academic Press, pp. 524-529, 2007.
- Dunn AJ, Swiergiel AH. The role of corticotropin-releasing factor and noradrenaline in stress-related responses, and the inter-relationships between the two systems. *Eur J Pharmacol* 583:186-193, 2008.
- Kino T., Charmandari E, Chrousos G.P. Disorders of the Hypothalamic-Pituitary-Adrenocortical System. In *Handbook of Neuroendocrinology* (Fink G., Pfaff DW, Levine J. eds). USA, NY: Academic Press, pp. 639-657, 2012.
- Koubassov RV, Barachevsky YE, Lupachev VV. Adrenocorticotrophic hormone and cortisol secretion changes among the law enforcement personnel during the mission to the area of armed conflicts. *Int J Biomed* 4:76-78, 2014.
- Selye H. *Stress without distress*. Philadelphia, USA: Lippincott, p. 171, 1974.
- Thrivikraman K.V, Nemeroff CB, Plotsky PM.

Sensitivity to glucocorticoid-mediated fast-feedback regulation of the hypothalamic-pituitary-adrenal axis is dependent upon stressor specific neurocircuitry. *Brain Res* 870:87-101, 2000.

15. Buckingham JC. Glucocorticoids, Role in Stress. In *Encyclopedia of Stress* (Fink G. ed). USA: Academic Press, pp. 210–217, 2007.

16. Morgan CA, Wang S, Rasmusson A, Hazlett G, Anderson G, Charney, DS. Relationship among plasma cortisol, catecholamines, neuropeptide Y, and human performance during exposure to uncontrollable stress. *Psychosom Med* 63:412-422, 2001.

17. Yehuda R. Current status of cortisol findings in post-traumatic stress disorder. *Psychiatr Clin North Am* 25:341-368, 2002.

18. Bremner JD, Vythilingam M, Vermetten E, Adil J, Khan S, Nazeer A, Afzal N, McGlashan T, Elzinga B, Anderson GM, Heninger G, Southwick SM, Charney DS. Cortisol response to a cognitive stress challenge in

posttraumatic stress disorder (PTSD) related to childhood abuse. *Psychoneuroendocrinology* 28:733-750, 2003.

19. Auxemery Y. Etiopathogenic perspectives on chronic psycho traumatic and chronic psychotic symptoms: The hypothesis of a hyperdopaminergic endophenotype of PTSD. *Med Hypotheses*. 79:667-672, 2012.

20. Emerson AJ, Kappenman DP, Ronan PJ, Renner KJ, Summers CH. Stress induces rapid changes in serotonergic activity: restraint and exertion. *Behav Brain Res* 111:83-92.

21. Eiden LE. Neuropeptide–Catecholamine Interactions in Stress. In: *A New Era of Catecholamines in the Laboratory and Clinic*. USA, Elsevier Inc. 68:399-404, 2013.

22. Murburg MM, McFall ME, Ko GN, Veith RC. Sympathoadrenal response to combat-related versus combat-unrelated stressors in combat veterans with post-traumatic stress disorder (PTSD) and controls. *Biol Psychiatry* 25:A33-A34, 1989.