A BIONIC APPROACH TO CARDIOVASCULAR REGULATION: BIONIC ARTERIAL BAROREFLEX SYSTEM

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Abstract- A bionic system is an artificial device, integrated into natural human physiology by communicating with native regulatory system. This can functionally operate as if it were a part of the body. Bionic systems can be realized only with the knowledge of detailed characteristics of native system. We made use of “white-noise approach” and have succeeded in functionally identifying the native arterial baroreflex. Using thus identified characteristics, we developed a bionic baroreflex system. Animal experiments in rats revealed that the bionic baroreflex system can stabilize pressure against hypotensive stimuli such as head-up tilt even without the native baroreflex system.

Keywords- White noise approach, Transfer function, Bionic baroreflex system

I. INTRODUCTION

A bionic system is an artificial device, integrated into natural human physiological systems by communicating with the native regulatory system, especially through the nervous system. Bionic systems can functionally operate as if they were a part of the body. These bionic systems open up a new therapeutic strategy against various cardiovascular diseases by restoring the lost function, or by modifying the abnormally functioning regulatory system.

Besides the methods to interface with the nervous system, development of bionic systems requires the knowledge of detailed characteristics of native system. Unlike artificial systems, native systems have a number of factors contributing to the complexity such as history-dependence, feedback nature, multiplicity and interaction of inputs, nonlinearity, and distributed nature. To overcome these, we made use of “white-noise approach” and have succeeded in functionally identifying the native arterial baroreflex with the precision one can reproduce native system artificially [1, 2, 3]. Using thus identified characteristics, we developed a bionic baroreflex system (Fig. 1) [4, 5] and examined if the bionic baroreflex system can stabilize pressure against hypotensive stimuli even in animals without functional baroreflex system.

II. METHODOLOGY

A. Functional identification with white-noise approach

In anesthetized rats with halothane, both carotid sinuses were vascularly isolated. Aortic depressor nerves and vagal nerves were cut bilaterally. We connected a high-fidelity piston pump to control carotid sinus pressure (CSP) as needed. We recorded CSP, cardiac sympathetic nerve activity (SNA), and systemic arterial pressure (SAP) simultaneously at sampling frequency of 100 Hz and resolution of 12 bits for 20 minutes while randomly perturbing sinus pressure. Nerve activity was quantified by smoothing the fully rectified signals. These signals were saved on hard disk for subsequent analysis. Transfer function (H_total) of total native baroreflex system was obtained.

Similarly we recorded blood pressure while stimulating sympathetic nerves at the celiac ganglia randomly. We
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obtained transfer function ($H_{effect}$) of the effector of native baroreflex. This transfer function strikingly resembles that from SNA and SAP obtained with CSP perturbation.

From these transfer functions, we calculated the dynamic characteristics ($H_{bionic}$), by dividing $H_{total}$ by $H_{effect}$ that are necessary for the bionic baroreflex system.

B. Functional replacement of baroreflex system with bionic approach

We used the same preparation for examining the bionic baroreflex performance as that described in the previous section. Although both carotid sinuses were isolated, pressure regulation under the operation of closed-loop baroreflex can be simulated by equating the sinus pressure to the systemic pressure with the piston pump. We abolished the baroreflex by keeping the sinus pressure constant. The bionic baroreflex was realized by stimulating the celiac ganglia according to the stimulation command. The stimulation command was calculated by convolving the instantaneous sinus pressure with the impulse response of the $H_{bionic}$. We examined the performance of bionic baroreflex system by imposing head-up tilt on these rats.

III. RESULTS

A. Functional identification with white-noise approach

The left panel of Fig. 2 illustrates an example of time course of random CSP perturbation and SAP changes in response to CSP. Careful observation reveals that SAP decreased when CSP increased though SAP was unresponsive to rapid changes in CSP. Transfer function $H_{total}$ showed low-pass characteristics with the corner frequency of ~0.1 Hz. Transfer function $H_{effect}$ similarly showed low-pass characteristics. The resultant $H_{bionic}$ showed almost all-pass characteristics with some derivative property in the high frequencies (Fig. 2, right).

B. Functional replacement of baroreflex system with bionic approach

As shown in Fig. 3, systemic pressure decreases dramatically in 10 seconds ($52 \pm 5$ mmHg, n=16) by the abrupt head-up tilt in rats with no functional baroreflex system. When we activates the bionic baroreflex system, thanks to the abrupt onset of celiac stimulation and increase in stimulation frequency, pressure considerably recovered in 10 seconds and decreased significantly less ($15 \pm 6$ mmHg). The time course and the amplitude of the hypotension with a bionic baroreflex were indistinguishable with a native one.

IV. DISCUSSION

Bionic medicine is a novel therapeutic strategy to treat various diseases by intervening native physiological system with bionic devices. Because the abnormality in cardiovascular regulation contributes to the maintenance of different cardiovascular diseases, this approach is likely to have a large number of applications in cardiology.

We have shown in this study that the bionic approach is feasible in treating orthostatic hypotension occurring as a result of central baroreflex failure. This system became possible with the detailed identification of the native system. In identifying the system we avoided to cut the system into pieces; we treated the system as a whole integrated one. This is in contrast with recent reductionist approaches in medicine and biology. We believe that this integrated approach is essential in developing a number of novel therapeutic devices to save patients suffering from various diseases.
V. CONCLUSION

By functionally identifying the arterial baroreflex system and by replacing it with the bionic system, we succeeded in reproducing the static as well as dynamic characteristics of the native baroreflex by an artificial device. The bionic medicine will pioneer the new therapeutic strategy not only in cardiology but also in various fields of medicine.

REFERENCES


