

## **Animal-Assisted Therapy in Patients Hospitalized With Heart Failure**

Kathie M. Cole, Anna Gawlinski, Neil Steers and Jenny Kotlerman

Am J Crit Care 2007;16:575-585  
© 2007 American Association of Critical-Care Nurses  
Published online <http://www.ajcconline.org>

Personal use only. For copyright permission information:  
[http://ajcc.aacnjournals.org/cgi/external\\_ref?link\\_type=PERMISSIONDIRECT](http://ajcc.aacnjournals.org/cgi/external_ref?link_type=PERMISSIONDIRECT)

### **Subscription Information**

<http://ajcc.aacnjournals.org/subscriptions/>

### **Information for authors**

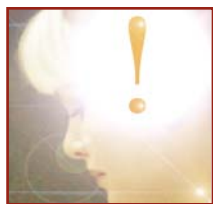
<http://ajcc.aacnjournals.org/misc/ifora.xhtml>

### **Submit a manuscript**

<http://www.editorialmanager.com/ajcc>

### **Email alerts**

<http://ajcc.aacnjournals.org/subscriptions/etoc.xhtml>



# ANIMAL-ASSISTED THERAPY IN PATIENTS HOSPITALIZED WITH HEART FAILURE

By Kathie M. Cole, RN, MN, CCRN, Anna Gawlinski, RN, DNSc, Neil Steers, PhD, and Jenny Kotlerman, MS

**CE** 1.0 Hour

### Notice to CE enrollees:

A closed-book, multiple-choice examination following this article tests your understanding of the following objectives:

1. Identify the physiological findings associated with advanced heart failure.
2. Discuss the overall effects of animal-assisted therapy on cardiopulmonary pressures, neurohormonal levels, and anxiety in advanced heart failure patients who participated in this study.
3. Describe the indications for further research in animal-assisted therapy that this study identifies.

To read this article and take the CE test online, visit [www.ajconline.org](http://www.ajconline.org) and click "CE Articles in This Issue." No CE test fee for AACN members.

**EBR**

Evidence-Based Review on pp 587-588.

**Background** Animal-assisted therapy improves physiological and psychosocial variables in healthy and hypertensive patients.

**Objectives** To determine whether a 12-minute hospital visit with a therapy dog improves hemodynamic measures, lowers neurohormone levels, and decreases state anxiety in patients with advanced heart failure.

**Methods** A 3-group randomized repeated-measures experimental design was used in 76 adults. Longitudinal analysis was used to model differences among the 3 groups at 3 times. One group received a 12-minute visit from a volunteer with a therapy dog; another group, a 12-minute visit from a volunteer; and the control group, usual care. Data were collected at baseline, at 8 minutes, and at 16 minutes.

**Results** Compared with controls, the volunteer-dog group had significantly greater decreases in systolic pulmonary artery pressure during (-4.32 mm Hg,  $P = .03$ ) and after (-5.78 mm Hg,  $P = .001$ ) and in pulmonary capillary wedge pressure during (-2.74 mm Hg,  $P = .01$ ) and after (-4.31 mm Hg,  $P = .001$ ) the intervention. Compared with the volunteer-only group, the volunteer-dog group had significantly greater decreases in epinephrine levels during (-15.86 pg/mL,  $P = .04$ ) and after (-17.54 pg/mL,  $P = .04$ ) and in norepinephrine levels during (-232.36 pg/mL,  $P = .02$ ) and after (-240.14 pg/mL,  $P = .02$ ) the intervention. After the intervention, the volunteer-dog group had the greatest decrease from baseline in state anxiety sum score compared with the volunteer-only (-6.65 units,  $P = .002$ ) and the control groups (-9.13 units,  $P < .001$ ).

**Conclusions** Animal-assisted therapy improves cardiopulmonary pressures, neurohormone levels, and anxiety in patients hospitalized with heart failure. (*American Journal of Critical Care*. 2007;16:575-588)

**H**earth failure is among the most common diagnoses in hospitalized adults in the United States. It is responsible for nearly 1 million hospitalizations annually, with estimated related healthcare costs of \$27.9 billion.<sup>1</sup> Hospitalization for heart failure is associated with a poor prognosis for patients, and readmission rates within 6 months are close to 50%.<sup>2</sup>

Heart failure induces a number of neurohormonal changes, including activation of the sympathetic nervous system, activation of the renin-angiotensin system, and reduction in activity of the parasympathetic nervous system.<sup>3</sup> Increased levels of catecholamines, such as epinephrine and norepinephrine, are hallmarks of the deleterious neuroendocrine cascade that occurs in patients with advanced heart failure. Stress-induced increases in the epinephrine level may facilitate release of norepinephrine.<sup>4</sup>

Although these changes may result in a short-term increase in cardiac output toward normal, chronic neurohormonal activation is harmful and contributes to progression of heart failure.<sup>5-7</sup> Chronic neurohormonal activation has led to the use of medications such as neurohormonal antagonists, including angiotensin-converting enzyme inhibitors, aldosterone antagonists, and  $\beta$ -adrenergic receptor antagonists, to treat heart failure.<sup>8</sup>

Although advances in medication therapy have improved outcomes of patients with heart failure, medication regimens have an unintended consequence of making polypharmacy a central component of the management of heart failure. Little is known about the hemodynamic and neurohormonal effects of adding adjunctive and complementary therapies to pharmacological management of advanced heart failure. Animal-assisted therapy (AAT) is an adjunctive therapy that could benefit patients with heart failure.

#### About the Authors

**Kathie M. Cole** is a clinical nurse III in the cardiac care unit, **Anna Gawlinski** is the director of evidence-based practice and an adjunct professor, and **Jenny Kotlerman** is a statistician at the Medical Center and School of Nursing, University of California, Los Angeles. **Neil Steers** is an adjunct assistant professor at the David Geffen School of Medicine, University of California, Los Angeles.

**Corresponding author:** Kathie M. Cole, RN, MN (CNIII), UCLA Medical Center, 4W CCU (Rm 46220), Los Angeles, CA 90095 (e-mail: nskmc@mednet.ucla.edu).

#### Review of the Literature

In AAT, the bond between humans and animals is an integral part of a patient's treatment.<sup>9</sup> Physiological variables change during short-term (2-12 minutes) interactions with animals and with pet ownership.<sup>10-13</sup> In several studies<sup>10,12-14</sup> with participants with normal or high blood pressure, interaction with animals yielded decreases in blood pressure and heart rate and an increase in peripheral skin temperature. Psychosocial and emotional benefits also have been the focus of studies of brief exposures of AAT of 10 to 30 minutes.<sup>15,16</sup> Psychosocial benefits include decreases in anxiety, isolation, and fear of procedures and improvements in social interaction, social support, communication, sensory stimulation, and happiness.<sup>15-21</sup>

In one study,<sup>22</sup> patients who were pet owners with long-term animal exposure had lower blood pressure, heart rate, and plasma renin activity in response to mental stressors (mathematical subtraction, speech) than did patients who were not pet owners. In patients who survived myocardial infarction, the risk for cardiovascular disease, morbidity, and mortality 1 year after the infarction was lower in those who were pet owners than in those who were not.<sup>23-25</sup> In the Cardiac Arrhythmia Suppression Trial, dog ownership was a significant independent predictor of survival in patients 1 year after acute myocardial infarction.<sup>23</sup> These data support the hypothesis that excess activity of the sympathetic nervous system due to both physiological and psychological stress can be reduced by AAT.

Patients with advanced heart failure are threatened by many physiological and psychological stressors.<sup>26,27</sup> Physiological stressors include the hallmark activation of the neuroendocrine cascade, most likely triggered by excitation of the sympathetic nervous system.<sup>3,28</sup> Chronic neurohormonal activation leads to ventricular remodeling.<sup>27</sup> Added to the physiological stress of heart failure is the psychological stress of living with a chronic, life-threatening illness and frequent hospitalizations. The presence of animals, and interaction with animals, decreases physiological indices such as heart rate and blood pressure and

Dog ownership is a significant, independent predictor of survival 1 year after a myocardial infarction.

improves psychosocial variables (eg, reduces anxiety) in both patients and healthy persons.<sup>15,22,29,30</sup> In a patient with heart failure, the presence of a non-threatening stimulus such as a dog could relax the patient by lowering the patient's state of arousal and reduce neurohormonal activation caused by overactivity of the sympathetic nervous system.<sup>31-33</sup>

Some patients may be indifferent to animals or may perceive animals as a source of stress.<sup>34</sup> In a study<sup>35</sup> of 58 hospitalized geriatric psychiatry patients who were randomly assigned to a pet therapy group or an exercise control group for 1 hour per day for 5 consecutive days, neither intervention resulted in significant differences in behavior or affect.<sup>35</sup> Similarly, in 33 male college students given mental stress tests with and without a dog present, no significant differences in dependent measures such as blood pressure and heart rate occurred between the 2 groups.<sup>36,37</sup>

Possible hazards associated with AAT include zoonotic infections (ie, infections that can be passed from animals to humans).<sup>34</sup> Effective strategies to prevent transmission of zoonotic infections include good hand washing and developing guidelines that include criteria for patient and animal suitability, infection control practices, and institutional policies.<sup>34,38</sup>

After the initiation of an AAT program, no zoonotic infections were reported in 3281 dog visits to 1690 hospitalized patients during a 5-year period.<sup>39</sup> Similarly, after the introduction of an AAT program, a children's hospital reported no increase in the rate of zoonotic infections or any other adverse incidences in the first 2-year period.<sup>40</sup>

Despite the applicability of AAT to hospitalized patients with heart failure, no randomized controlled trials of the effects of this therapy have been done. The purpose of this study was to determine if AAT could reduce the manifestations of physiological and psychological stress in patients with advanced heart failure. Specifically, we tested whether hospitalized patients with advanced heart failure who received AAT had improved hemodynamic measures, lower neurohormone levels, and decreased anxiety compared with patients visited by a volunteer only and a control group of patients who received usual care at rest.

## Methods

### Design

A 3-group (volunteer-dog team, volunteer only, and control) randomized, repeated-measures experimental design was used to determine the effect of AAT on multiple dependent variables. Dependent variables were (1) blood pressure, (2) heart rate, (3) pulmonary artery pressure (PAP), (4) pulmonary

capillary wedge pressure (PCWP), (5) right atrial pressure (RAP), (6) cardiac index, (7) systemic vascular resistance (SVR), (8) epinephrine level, (9) norepinephrine level, and (10) state anxiety.

To control for the effect of the volunteer part of the team, we designed the study to include a group visited by a volunteer only and a control group. The experimental group received the AAT, which consisted of a 12-minute visit from a volunteer and dog per standard AAT protocol (see "Data Collection Procedures" section). The AAT protocol has been used at the University of California–Los Angeles Medical Center since 1994.<sup>41</sup> The patients in the volunteer-only group received a 12-minute visit from a volunteer who was unknown to the patients. The patients in the control group received usual care (at rest).

### Sample and Setting

After approval from the institutional review board, 76 patients with a diagnosis of advanced heart failure admitted to the cardiac care unit or the cardiac observation unit were recruited for the study. Patients who met the selection criteria, agreed to participate, and signed the informed consent were randomized into 1 of 3 groups (volunteer and dog visit, visit by volunteer only, control). Volunteers also signed an informed consent as requested by the institutional review board.

Criteria for selection of participants included (1) advanced heart failure (including systolic and diastolic left ventricular dysfunction) requiring medical management with an indwelling pulmonary artery catheter; (2) age between 18 and 80 years; (3) ability to read, write, and speak English; (4) mental status alert and oriented to person, place, and time; and (5) SVR greater than  $1200 \text{ dyne} \cdot \text{sec} \cdot \text{cm}^{-5}$  at least once within 12 hours from the start of data collection. Exclusion criteria included (1) SVR less than  $1200 \text{ dyne} \cdot \text{sec} \cdot \text{cm}^{-5}$ ; (2) allergies to dogs; (3) immunosuppression, defined as a white blood cell count of less than 4500 cells/ $\mu\text{L}$ ; (4) infection, indicated by an elevated white blood cell count greater than 11 000 cells/ $\mu\text{L}$ ; (5) body temperature greater than  $38^\circ\text{C}$ ; and (6) decreased level of consciousness.

The presence of and interaction with animals decreases heart rate and blood pressure and improves anxiety.

### Data Collection Procedures

Patients were randomly assigned to 1 of 3 groups by using a table of random numbers. Group assignment determined the type of visit. Patients randomly assigned to the experimental group received a visit from a volunteer and a dog. The type of dog breed was not controlled for; 14 dogs of 10 various breeds were used. The 14 dogs included 1 extra-large dog, 6 large dogs, 5 medium dogs, and 2 small dogs. Each visit was conducted according to the guidelines taught during the volunteer and dog orientation: (1) volunteer introduces self and dog, (2) patient washes his or her hands before the visit, (3) dog lies on the bed with its head within 0.6 m (2 ft) of the patient's head on a clean sheet used as a barrier to the patient's bed, (4) patients may pet the dog and talk to the dog and volunteer, and (5) patient washes his or her hands after the visit. No attempt was made to control the content of the conversation during the visit. The visit lasted for 12 minutes. After the visit, an instant self-developing photograph was taken of the patient with the dog and given to the patient.

Patients randomly assigned to the volunteer-only group received a 12-minute visit from a volunteer.

The volunteer introduced himself or herself, sat in a chair approximately 1.2 m (4 ft) from the patient's head, and let the patient know that the visit would last for 12 minutes if the patient was up to it. No attempt was made to control for the volunteer's usual conversation during the visit. No patient requested to end any earlier than 12 minutes. Patients randomly assigned to the control group were asked to lie quietly without talking unless they had a specific need or request. For all groups, a sign was placed on the

patient's door or curtain asking everyone to please not interrupt the visit. Nurses assigned to patients to provide care were asked not to interrupt during the 12-minute interaction and data collection, unless an emergency occurred. Volunteers participating in the volunteer-dog teams were used for the volunteer-only group as much as possible to minimize any influence of a volunteer's personality on the results.

For all groups, data were collected at baseline immediately before the visit, 8 minutes after the intervention started, and at 16 minutes, which was 4 minutes after the end of the visit. These times were chosen because they most likely correspond to the maximal relaxation effect for a patient<sup>18,30</sup> and thus were times when a difference from baseline levels

was most likely to be detected. For all data collection, patients were recumbent with the head of bed elevated 45° from horizontal.

The physiological variables (blood pressure, heart rate, PAP, PCWP, RAP, cardiac index, SVR, epinephrine level, and norepinephrine level) were assessed at all 3 times for all groups. In order to maximize a "steady state" period, neither dosages of intravenous medications nor patients' positions were changed in the 15 minutes before data collection or during data collection. Blood samples for measurement of plasma levels of epinephrine and norepinephrine were obtained at all 3 times for all groups. Anxiety was measured twice for all groups: at baseline and at 16 minutes. Only 2 measures of anxiety were used to avoid sensitizing patients to the instrument.

All data were collected by the coinvestigator (K.M.C.) or research assistants, who were critical care nurses with expertise in all areas of hemodynamic monitoring. Data collectors did not speak to the patients during the measurement of outcome variables and the intervention. Interrater reliability ( $r=0.94$ ) was established among research assistants by providing instruction that included training on how to measure all the dependent variables and completion of a test after the training on accurate measurement of hemodynamic variables. The entire data collection time per patient was approximately 1 hour.

### Measures

#### Hemodynamic Parameters

Heart rate is defined as the number of beats per minute and was measured by using a bedside monitor (model M1176A, Phillips Medical Systems, Andover, Massachusetts). Blood pressure was measured noninvasively and automatically by using a bedside monitor (model M1006B, Phillips Medical Systems). The reliability and validity of the noninvasive automatic measurements of blood pressure have been established, and the accuracy of these measurements compared with intra-arterial measurements is high, with observed mean errors of +0.8% for systolic blood pressure and +1.7% for diastolic blood pressure.<sup>42</sup>

Cardiac index is a measure of flow. It is calculated as cardiac output in liters per minute divided by body surface area in meters squared. Cardiac output was measured by using the thermodilution technique: 3 injections with 10 mL of iced injectate per injection. Results were averaged to obtain cardiac output in liters per minute. SVR is total peripheral resistance of the vascular system and reflects the degree of vascular constriction. SVR was calculated from cardiac output by using the following formula:

The volunteer-dog team visit resulted in decreases in pulmonary artery and wedge pressures and state anxiety levels.

([mean arterial pressure - RAP] x 80) divided by the cardiac output. Reliability of measures was ensured by leveling, zeroing, and calibrating instruments according to the manufacturers' specifications at the start of the study period. Accuracy of cardiac output obtained via the thermodilution method compared with output obtained via the direct Fick and dye dilution methods has been established ( $r = 0.91-0.97$ ).<sup>43</sup>

### Neurohormone Levels

Plasma levels of catecholamines (epinephrine and norepinephrine) were measured by using standard laboratory procedures (high-pressure liquid chromatography) and electromechanical detection. Because position can affect catecholamine levels, all patients were supine with the head of the bed elevated 45° when blood samples were obtained, and all had been in that position for 15 minutes before the first blood sample was taken. For measurements of catecholamine levels, 7 mL of blood was withdrawn from the proximal port of the pulmonary artery catheter after in-line withdrawal of an appropriate amount of blood for discard. Blood samples were placed in chilled heparinized tubes, put on ice, and immediately transported to the laboratory for processing. Reliability and validity of the measurement method have been well established. The interassay coefficient of variation was 8% for epinephrine and 5% for norepinephrine.<sup>44</sup>

### Anxiety

The Spielberger State-Trait Anxiety Inventory, form Y-I, was used to measure state anxiety, defined as a transitory emotional state or condition of the human individual, characterized by subjective, consciously perceived feelings of tension and apprehension and heightened arousal of the autonomic nervous system.<sup>45</sup> The State Anxiety Inventory is a self-report questionnaire that asks participants about their feelings of anxiety "right now." It consists of 20 statements such as "I feel at ease" and "I feel worried." Our patients rated themselves on a 4-point scale of increasing intensity. After reverse-coding necessary items, we computed the scale score as the sum of the 20 items. The instrument has well-established reliability (Cronbach  $\alpha$  .83-.92), validity (content, concurrent, and construct), and sensitivity for distinguishing changes in anxiety during periods as short as 10 to 30 minutes.<sup>45</sup>

### Data Analysis

Descriptive statistics were used for all dependent variables. Means and standard deviations were calculated for continuous variables. Differences among

the groups at baseline were examined by using 1-way analysis of variance. Frequencies and percentages were determined for categorical variables. The differences among the groups at baseline were determined by using the  $\chi^2$  statistic.

For each dependent variable, a mixed model was used to test for differences among the 3 groups at the 3 time points. All mixed model analyses were performed by using SAS statistical software (SAS Institute Inc, Cary, North Carolina). In each model, adjusted differences in the dependent variable from baseline to during the intervention (baseline-during differences) and from baseline to after the intervention (baseline-after differences) were estimated for all pairs of groups. For example, to compare the volunteer and dog team with the control group in the differences between baseline and during the intervention measurements, the estimated differences from baseline to during were calculated for each group and then subtracted by using the Estimate function in Proc Mixed. *P* values for these comparisons were computed. Each model was adjusted for the baseline measure of its dependent variable, age, sex, New York Heart Association classification, ejection fraction, dog ownership (never, in the past, present), having a live-in partner or a spouse, and current smoking status.

## Results

### Characteristics of the Sample

The 76 patients who completed the study had advanced heart failure as indicated by depressed left ventricular ejection fraction (mean, 23.4%; SD, 10.80%), impaired functional ability (New York Heart Association functional class III and IV,  $n = 72$ ), vasoconstriction (SVR, mean, 1372 dyne  $\cdot$  sec  $\cdot$  cm<sup>-5</sup>; SD, 365), and elevated plasma level of norepinephrine (mean, 1008 pg/mL; SD, 704; to convert to picomoles per liter, multiply by 5.911). Mean age of patients in the sample was 57 years (SD, 12.28). A total of 75% were men.

Heart failure was most often due to nonischemic cardiomyopathy (44 patients, 58%). Medications affecting cardiopulmonary status were not controlled for; however, intravenous infusions of drugs were not manipulated and oral medications were not given 1 hour before or during the intervention. The number and types of medications used did not differ significantly between groups; patients were given diuretics (83%), vasoactive medications (82%), inotropic agents (55%), angiotensin-converting enzyme

Animal-assisted therapy reduced epinephrine and norepinephrine levels, suggesting changes in activation of the autonomic nervous system.

**Table 1**  
Comparison of baseline demographic and clinical characteristics of the sample between volunteer-dog, volunteer-only, and control groups

Variable	Groups			P
	Volunteer-dog (n = 26)	Volunteer-only (n = 25)	Control (n = 25)	
	Mean (SD)	Mean (SD)	Mean (SD)	
Left ventricular ejection fraction, %	23.73 (12.26)	22.14 (9.76)	24.34 (11.04)	.77
Heart rate, beats/min	86.77 (18.83)	83.08 (13.87)	81.68 (15.45)	.50
Systolic blood pressure, mm Hg	98.85 (11.89)	100.68 (15.86)	99.56 (11.44)	.88
Diastolic blood pressure, mm Hg	60.69 (12.43)	61.68 (10.05)	59.04 (9.98)	.69
Mean arterial pressure, mm Hg	73.10 (10.43)	72.36 (11.06)	72.32 (9.18)	.95
Systolic pulmonary artery pressure, mm Hg	44.12 (14.23)	39.12 (10.54)	38.20 (11.19)	.18
Diastolic pulmonary artery pressure, mm Hg	20.96 (6.37)	19.52 (5.09)	19.64 (5.79)	.61
Pulmonary capillary wedge pressure, mm Hg	21.11 (6.51)	17.14 (5.79)	16.36 (4.74)	.02
Pulmonary capillary wedge pressure, mm Hg (without 2 outliers)	19.65 (5.11)	17.14 (5.79)	16.36 (4.74)	.15
Right atrial pressure, mm Hg	8.62 (5.12)	7.16 (3.88)	6.72 (3.63)	.26
Cardiac output, L/min	3.82 (0.92)	4.10 (0.79)	4.04 (0.66)	.41
Cardiac index <sup>a</sup>	2.02 (0.49)	2.14 (0.42)	2.13 (0.35)	.52
Systemic vascular resistance, dyne · sec · cm <sup>-5</sup>	1428.62 (422.90)	1361.36 (394.22)	1348.96 (287.45)	.71
Plasma norepinephrine, pg/mL <sup>b</sup>	1098.23 (733.93)	927.65 (475.46)	923.43 (651.43)	.58
Plasma epinephrine, pg/mL <sup>c</sup>	84.45 (59.54)	58.27 (42.21)	52.68 (44.79)	.10
Anxiety, sum score in units	44.35 (15.32)	37.83 (12.21)	40.04 (12.70)	.23
Age, y	56.27 (14.36)	56.12 (12.56)	58.52 (9.73)	.74
	No. (%)	No. (%)	No. (%)	P
Men	17 (65)	22 (88)	18 (72)	.16
New York Heart Association class IV	17 (65)	18 (72)	15 (60)	.67
20% Left ventricular ejection fraction	8 (31)	8 (32)	7 (28)	.95
Cause of heart failure				
Ischemic	8 (31)	12 (48)	12 (48)	.35
Current smoker	4 (15)	3 (12)	1 (4)	.40
Living with a spouse (or significant other)	20(77)	15 (60)	17 (68)	.43
Current dog owner	19 (73)	14 (56)	11 (44)	.10
Previous dog owner	24 (92)	24 (96)	22 (88)	.11

<sup>a</sup> Calculated as cardiac output in liters divided by body surface area in meters squared.

<sup>b</sup> To convert to picomoles per liter, multiply by 5.911.

<sup>c</sup> To convert to picomoles per liter, multiply by 5.459.

inhibitors (62%), and  $\beta$ -blockers (38%). A total of 58% of the patients were current dog owners. The only difference among the groups at baseline was PCWP. It was significantly higher in the patients who received a visit from the volunteer-dog team than in the other 2 groups ( $P = .02$ , Table 1).

#### Mixed Models

Adjusted mean baseline-during differences for the 3 groups are presented in Table 2. Several differences

in dependent variables were detected between the volunteer-dog group and the control group. Compared with the control group, patients in the volunteer-dog group had significantly greater decreases in systolic PAP (mean difference,  $-4.32$  mm Hg;  $P = .03$ ) and PCWP (mean difference,  $-2.74$  mm Hg;  $P = .01$ ).

Similar baseline-during differences were detected between the volunteer-dog group and the volunteer-only group (Table 2). Compared with the volunteer-only group, the volunteer-dog group had

**Table 2**  
Differences between groups in decreases in each dependent variable from baseline to during the intervention

Variable	Groups								
	Volunteer-dog vs control <sup>a</sup>			Volunteer-dog vs volunteer-only <sup>b</sup>			Volunteer-only vs control <sup>c</sup>		
	Adjusted mean difference (SD), <i>P</i>								
Heart rate, beats/min	-1.35	(1.84)	.47	-3.55	(1.84)	.06	+0.79	(1.35)	.56
Respiratory rate, respirations/min	-0.19	(0.64)	.76	-1.03	(0.64)	.11	+0.36	(0.59)	.54
Systolic blood pressure, mm Hg	+1.85	(2.30)	.42	-0.71	(2.30)	.76	-0.04	(2.25)	.99
Diastolic blood pressure, mm Hg	-0.42	(2.88)	.84	+0.18	(2.88)	.95	-0.88	(2.82)	.31
Mean arterial pressure, mm Hg	-1.23	(2.27)	.59	-0.69	(2.27)	.76	-0.59	(2.19)	.79
Systolic pulmonary artery pressure, mm Hg	-4.32	(1.99) <sup>d</sup>	.03	-5.48	(1.99) <sup>d</sup>	.01	+0.13	(1.47)	.93
Diastolic pulmonary artery pressure, mm Hg	-0.58	(1.39)	.67	-2.78	(1.39) <sup>d</sup>	.05	+1.19	(0.98)	.23
Pulmonary capillary wedge pressure, mm Hg	-2.74	(1.04) <sup>d</sup>	.01	-3.30	(1.04) <sup>d</sup>	.003	+0.21	(0.97)	.83
Pulmonary capillary wedge pressure, mm Hg (without 2 outliers)	-3.11	(0.06) <sup>d</sup>	.005	-3.70	(1.06) <sup>d</sup>	.001	+0.58	(0.99)	.56
Right atrial pressure, mm Hg	-0.27	(0.70)	.70	-1.87	(0.70) <sup>d</sup>	.01	+0.91	(0.57)	.11
Cardiac output, L/min	+0.14	(0.17)	.41	+0.16	(0.17)	.36	-0.32	(0.20)	.11
Cardiac index <sup>e</sup>	+0.11	(0.08)	.19	+0.09	(0.08)	.27	-0.11	(0.10)	.27
Systemic vascular resistance, dyne · sec · cm <sup>-5</sup>	+11.9	(85.3)	.89	-18.28	(85.27)	.83	+87.34	(91.90)	.35
Norepinephrine, pg/mL <sup>f</sup>	-47.44	(96.61)	.63	-232.36	(95) <sup>d</sup>	.02	-7.78	(93)	.93
Epinephrine, pg/mL <sup>g</sup>	-2.80	(8.18)	.73	-15.86	(7.82) <sup>d</sup>	.04	-1.68	(9.85)	.87

<sup>a</sup> Adjusted mean difference = mean decreases in the volunteer-dog group minus mean decreases in the control group. Negative values indicate greater decreases in the volunteer-dog group; positive values indicate greater decreases in the control group.

<sup>b</sup> Adjusted mean difference = mean decreases in the volunteer-dog group minus mean decreases in the volunteer-only group. Negative values indicate greater decreases in the volunteer-dog group; positive values indicate greater decreases in the volunteer-only group.

<sup>c</sup> Adjusted mean difference = mean decreases in the volunteer-only group minus mean decreases in the control group. Negative values indicate greater decreases in the volunteer-only group; positive values indicate greater decreases in the control group.

<sup>d</sup>  $P \leq .05$ .

<sup>e</sup> Calculated as cardiac output in liters divided by body surface area in meters squared.

<sup>f</sup> To convert to picomoles per liter, multiply by 5.911.

<sup>g</sup> To convert to picomoles per liter, multiply by 5.459.

significantly greater decreases in systolic PAP (mean difference, -5.48 mm Hg;  $P = .01$ ), PCWP (mean difference, -3.30 mm Hg;  $P = .003$ ), epinephrine levels (mean difference, -15.86 pg/mL;  $P = .04$ ; to convert to picomoles per liter, multiply by 5.459), and norepinephrine levels (mean difference, -232.36 pg/mL;  $P = .02$ ). In addition, patients in the volunteer-dog group had greater decreases in RAP (mean difference, -1.87 mm Hg;  $P = .01$ ) and diastolic PAP (mean difference, -2.78 mm Hg;  $P = .05$ ).

Adjusted mean baseline-after differences for the 3 groups are presented in Table 3. Two of the mean baseline-during differences between the volunteer-dog group and the control group also were found for mean baseline-after differences. Compared with the control group, the volunteer-dog group had significantly greater decreases in systolic PAP (mean difference, -5.78 mm Hg;  $P = .001$ ) and PCWP

(mean difference, -4.31 mm Hg;  $P = .001$ ). Decreases in epinephrine and norepinephrine were similar for the 2 groups. However, compared with the control group, the volunteer-dog group had greater reductions in state anxiety (mean difference, -9.13 units;  $P < .001$ ).

Most of the baseline-during differences between the volunteer-dog group and the volunteer-only group also were found for baseline-after differences (Table 3). Compared with the volunteer-only group, the volunteer-dog group had significantly greater decreases in systolic PAP (mean difference, -5.34 mm Hg;  $P = .002$ ), PCWP (mean difference, -3.10 mm Hg;  $P = .02$ ); epinephrine levels (mean difference, -17.54 pg/mL;  $P = .04$ ), and norepinephrine levels (mean difference, -240.29 pg/mL;  $P = .02$ ). However, decreases in RAP and diastolic PAP in the volunteer-dog group were not significantly greater than those in the volunteer group. In addition, the



**Table 3**

Differences between groups in decreases in each dependent variable from baseline to after the intervention

Variable	Groups								
	Volunteer-dog vs control <sup>a</sup>			Volunteer-dog vs volunteer-only <sup>b</sup>			Volunteer-only vs control <sup>c</sup>		
Adjusted mean difference (SD), P									
Heart rate, beats/min	-1.44	(1.49)	.34	-2.76	(1.49)	.07	+1.32	(1.50)	.38
Respiratory rate, respirations/min	-0.39	(0.67)	.56	-0.67	(0.66)	.32	-0.27	(0.67)	.68
Systolic blood pressure, mm Hg	-3.35	(2.46)	.18	-0.75	(2.46)	.76	-2.60	(2.49)	.30
Diastolic blood pressure, mm Hg	-4.38	(2.53)	.09	-2.70	(2.53)	.29	-1.68	(2.56)	.51
Mean arterial pressure, mm Hg	-3.36	(2.25)	.14	-1.28	(2.25)	.57	-2.08	(2.27)	.36
Systolic pulmonary artery pressure, mm Hg	-5.78	(1.73) <sup>d</sup>	.001	-5.34	(1.72) <sup>d</sup>	.002	-0.44	(1.74)	.80
Diastolic pulmonary artery pressure, mm Hg	-0.68	(1.38)	.63	-1.59	(1.38)	.25	+0.92	(1.39)	.51
Pulmonary capillary wedge pressure, mm Hg	-4.31	(1.30) <sup>d</sup>	.001	-3.10	(1.28) <sup>d</sup>	.02	-1.22	(1.25)	.34
Pulmonary capillary wedge pressure, mm Hg (without 2 outliers)	-4.29	(1.36) <sup>d</sup>	.003	-3.07	(1.34) <sup>d</sup>	.03	-1.22	(1.27)	.34
Right atrial pressure, mm Hg	+0.44	(0.55)	.43	-0.96	(0.55)	.09	+1.40	(0.56) <sup>d</sup>	.01
Cardiac output, L/min	+0.06	(0.18)	.74	-0.16	(0.18)	.38	+0.22	(0.18)	.23
Cardiac index <sup>e</sup>	+0.10	(0.10)	.31	-0.02	(0.10)	.86	+0.12	(0.10)	.24
Systemic vascular resistance, dyne · sec · cm <sup>-5</sup>	-73.30	(78.02)	.35	+69.06	(78.02)	.38	-142.36	(78.79)	.08
Norepinephrine, pg/mL <sup>f</sup>	-182.31	(102)	.08	-240.14	(100) <sup>d</sup>	.02	+57.83	(101)	.57
Epinephrine, pg/mL <sup>g</sup>	-12.56	(8.66)	.15	-17.54	(8.28) <sup>d</sup>	.04	+4.98	(8.46)	.56
Anxiety, sum score in units	-9.13	(2.10) <sup>d</sup>	<.001	-6.65	(2.13) <sup>d</sup>	.002	-2.48	(2.17)	.25

<sup>a</sup> Adjusted mean difference = mean decreases in the volunteer-dog group minus mean decreases in the control group. Negative values indicate greater decreases in the volunteer-dog group; positive values indicate greater decreases in the control group.

<sup>b</sup> Adjusted mean difference = mean decreases in the volunteer-dog group minus mean decreases in the volunteer-only group. Negative values indicate greater decreases in the volunteer-dog group; positive values indicate greater decreases in the volunteer-only group.

<sup>c</sup> Adjusted mean difference = mean decreases in the volunteer-only group minus mean decreases in the control group. Negative values indicate greater decreases in the volunteer-only group; positive values indicate greater decreases in the control group.

<sup>d</sup>  $P \leq .05$ .

<sup>e</sup> Calculated as cardiac output in liters divided by body surface area in meters squared.

<sup>f</sup> To convert to picomoles per liter, multiply by 5.911.

<sup>g</sup> To convert to picomoles per liter, multiply by 5.459.

volunteer-dog group had significantly greater decreases in state anxiety (mean difference, -6.65 units,  $P = .002$ ) than did the volunteer-only group.

Because PCWP was the only variable at baseline that differed significantly between the 3 groups, we explored this variable further. Using the box-plot method, we calculated the distance 1.5 times the interquartile range (ie, the distance between the 75th and 25th percentiles of the data). Any value more than this distance away from the median in either direction was considered an outlier. The volunteer-dog group had 2 outlier values. After eliminating these outlier values, we repeated the analysis for differences in PCWP between groups. The results showed similar differences between groups in PCWP with and without the outliers (Tables 2 and 3).

Therefore, the baseline-during and baseline-after values for systolic PAP and PCWP decreased significantly more in the volunteer-dog group than in the volunteer-only and control groups. Similarly, baseline-during and baseline-after values for epinephrine and norepinephrine levels decreased significantly more in the volunteer-dog group than in the volunteer-only group. The baseline-after decrease in state anxiety was greater in the volunteer-dog group than in the other 2 groups.

## Discussion

In this study, patients hospitalized with advanced heart failure who received a visit from a volunteer and dog had lower cardiopulmonary pressures, neurohormone levels, and anxiety levels than did patients visited by a volunteer only and patients

given usual care at rest (control group). Heart rate, blood pressure, cardiac index, and SVR were not significantly affected by the intervention.

In previous studies,<sup>10,22,46</sup> the presence of a dog, pet ownership, and AAT resulted in lower blood pressures in hypertensive and normotensive pet owners and in participants who did not own a pet. The mean blood pressure in these studies<sup>10,22,46</sup> ranged from a systolic pressure of 116 to 182 mm Hg and a diastolic pressure of 81 to 120 mm Hg. Several investigators<sup>25,31,33</sup> postulated that changes in blood pressure in pet owners were due to a decrease in sympathetic tone; however, the investigators did not measure neuroendocrine markers such as catecholamine levels.

The patients in our study did not have decreases in blood pressure similar to those reported in previous studies. One explanation is that changes in blood pressure in our patients may have been blunted by severe preexisting cardiac dysfunction. In addition, several of our patients received medications that could affect blood pressure response. A total of 62 patients (82%) were receiving vasoactive medications and 42 (55%) were receiving inotropic medications. Mean baseline blood pressures in our patients with advanced heart failure were low (mean systolic pressure, 99 mm Hg; mean diastolic pressure, 60 mm Hg) compared with normal and hypertensive populations in other animal interaction studies (mean systolic pressure, 144 mm Hg; mean diastolic pressure, 86 mm Hg).<sup>10,11,13,14,22</sup>

Despite evidence that AAT can lower blood pressure, the hemodynamic and neuroendocrine effects of this therapy are relatively unknown. To date, our study is the only experimental research in which the hemodynamic and neuroendocrine responses to AAT were examined. Hemodynamic effects of AAT included significant decreases in systolic PAP and PCWP during and after the intervention in patients visited by the volunteer and dog compared with the patients in the other 2 groups. Diastolic PAP and RAP were significantly lower during the intervention in the volunteer-dog group than in the volunteer-only group. Previous research<sup>47,48</sup> showed that the simple task of talking about a neutral topic can elevate cardiovascular reactivity, as indicated by increases in systolic blood pressure, diastolic blood pressure, and heart rate in patients with cardiovascular disorders.

Nemens and Woods<sup>49</sup> examined normal fluctuations in baseline hemodynamic pressures in a sample of cardiovascular patients and noted fluctuations of 5 mm Hg in systolic PAP. In our study, the mean decreases in systolic PAP from baseline to after the AAT intervention were approximately 6 mm Hg greater in the volunteer-dog group than in the control

group. Therefore, the reduction in systolic PAP in response to AAT in the hospitalized patients with heart failure in our group was clinically relevant.

Hemodynamic changes also were associated with changes in plasma levels of epinephrine and norepinephrine. The decreases in plasma levels of epinephrine and norepinephrine during and after the intervention were significantly greater in the volunteer-dog group than in the volunteer-only group.

Plasma levels of catecholamines reflect the balance between the release of the hormones from sympathetic nerve terminals and their reuptake and degradation.<sup>50</sup> Increased levels of catecholamines, such as epinephrine and norepinephrine, are hallmarks of the deleterious neuroendocrine cascade that occurs in patients with advanced heart failure. The inotropic effects of endogenous catecholamines are blunted in heart failure, and catecholamines have direct deleterious effects on myocardial cells, left ventricular function, and mortality.<sup>51,52</sup> Furthermore, when a person perceives fear or loss of control, brain neuropeptides activate catecholamines such as epinephrine and norepinephrine, which affect heart rate and blood pressure.<sup>53</sup>

In our study, the decreases in neurohormone levels during and after AAT suggest changes in the activation of the autonomic nervous system. AAT may affect neurohormone levels by altering the response of the autonomic nervous system to stimuli that are perceived as pleasantly meaningful and stimuli that the individual wishes to interact with from the environment.<sup>54</sup> Thus, patients receiving AAT may have been focused on the dog and not on other environmental stimuli, such as the volunteer, the nurse who collected data, and other sensory stimuli present at the time of the intervention.

In our study, the decrease in anxiety after the intervention was significantly greater in the volunteer-dog group than in the volunteer-only and the control groups. Similar findings have been observed in hospitalized psychiatric patients and in college students.<sup>15,17</sup> For example, Barker and Dawson<sup>15</sup> investigated the effects of AAT on 230 hospitalized psychiatric patients and compared a single 30-minute AAT session with a 30-minute therapeutic recreation session. A significant decrease in anxiety occurred in patients with mood disorders, psychotic disorders, and other disorders who received AAT sessions.

The significant decrease in anxiety in the volunteer-dog group suggests that AAT may provide patients a meaningful stimulus and modulate anxiety-producing situations that encompass hospitalization, severe illness, and uncertainty of outcomes. AAT may be effective because pets are a source of social support and

may buffer a person's reactivity to mental stressors.<sup>20,22,23</sup> AAT can provide a sense of comfort and safety during hospitalization, and it diverts attention away from the immediate stressors to a more pleasurable and calming interaction.<sup>15</sup>

After the intervention, the volunteer-only group had a significantly greater decrease in state anxiety than did the control group, who received usual care at rest. However, AAT resulted in the greatest decrease in state anxiety scores and was more effective than a visit from a volunteer only or leaving a patient at rest.

## Limitations

The study has several limitations. Although PCWP was the only variable that differed significantly between the 3 groups at baseline, the volunteer-dog group had higher observed means than did the other 2 groups for most of the dependent variables at baseline. Data collection occurred during a short period (12 minutes), and patients in the volunteer-dog group with higher baseline measures might have been treated more aggressively than patients in the other 2 groups. Although no medication changes occurred 1 hour before and during the intervention, the greatest decreases in PAP, PCWP, and neurohormones could have been influenced by ongoing intravenous drug infusions at the time. Thus, the study results should be viewed cautiously, and replication is needed.

The lack of changes in other hemodynamic parameters such as cardiac index and SVR may be due to the short exposure (12 minutes) of the patients to the AAT. It is not known whether longer exposures may result in additional hemodynamic benefits. In addition, fluid limitations precluded more frequent measurements of cardiac index. Finally, the study was conducted at a single facility.

## Conclusions

Our findings add to knowledge on AAT and indicate a new focus for study of interactions between physiological and psychological effects and therapeutic benefits of AAT. Although this preliminary study provides evidence of improved cardiopulmonary pressures, neurohormone levels, and anxiety in patients with heart failure who have AAT, a larger study is needed to sufficiently define the immediate and long-term improvement in these variables. Healthcare professionals can use AAT as an effective adjunctive treatment.

Further research is needed to evaluate the potential effect of AAT for achieving better management of symptoms (eg, dyspnea), improving satisfaction of

patients, and decreasing length of stay in the hospital. Other areas of investigation are needed to determine whether this safe, inexpensive adjunctive therapy may contribute to the long-term treatment of patients with advanced heart failure by decreasing morbidity and mortality, decreasing depression, and improving the quality of life in domains such as functional status and social support.

## ACKNOWLEDGMENTS

We acknowledge the following for their contributions: research professor Donna Vredavoe, PhD, and Priscilla Kehoe, RN, PhD, at the University of California, Los Angeles, School of Nursing for study consultation; Gregg C. Fonarow, MD, and Jan H. Tillisch, MD, for support and encouragement to conduct the study; the nursing staff of the cardiac care and cardiac observation units; research assistants Rosanne Giza, RN, BSN, Moira Kelly, RN, BSN, Magdelaine Panganiban, RN, MSN, Lance Patak, RN, BSN, and Patty Trapnell, RN; data entry reviewers Jhoanna Anuran-Torres, RN, MS, and Heather Saunders, RN, MS; Dr Kehoe for critical revision of the manuscript; the animal-assisted therapy program at the university, the People-Animal Connection, for providing volunteers and therapy dogs; and Lynda Oschin and the late Robert Oschin for providing the People-Animal Connection program philanthropy. We are indebted to the patients in the cardiac care and cardiac observation units; without them this study would not have been possible.

## FINANCIAL DISCLOSURES

This study was funded by grant 01061809 from the Pet Care Trust Foundation, Quilcene, Washington.

## eLetters

Now that you've read the article, create or contribute to an online discussion about this topic using eLetters. Just visit [www.ajconline.org](http://www.ajconline.org) and click "Respond to This Article" in either the full-text or PDF view of the article.

## SEE ALSO

To learn more about pet visiting and animal-assisted therapy, read chapter 4, "Family Pet Visiting and Animal-Assisted Therapy," by Nancy C. Molter, RN, MN, PhD, in *AACN Protocols for Practice: Creating Healing Environments* (Jones and Bartlett Publishers; 2007), pp 103-120. Available online at the AACN Bookstore: <http://www.aacn.org/creatinghealingenvironments>.

## REFERENCES

1. American Heart Association. *Heart Disease and Stroke Statistics: 2005 Update*. Dallas, TX: American Heart Association; 2005.
2. Krumholz HM, Parent EM, Tu N, et al. Readmission after hospitalization for congestive heart failure among Medicare beneficiaries. *Arch Intern Med*. 1997;157(1):99-104.
3. Schrier RW, Abraham WT. Hormones and hemodynamics in heart failure. *N Engl J Med*. 1999;341:577-585.
4. Esler M, Lambert G, Brunner-La Rocca HP, Vaddadi G, Kaye D. Sympathetic nerve activity and neurotransmitter release in humans: translation from pathophysiology into clinical practice. *Scand Physiol Soc*. 2003;177:275-284.
5. Bristow MR.  $\beta$ -Adrenergic receptor blockade in chronic heart failure. *Circulation*. 2000;101:558-569.
6. Piano MR, Bondmas M, Schwartz DW. The molecular and

- cellular pathophysiology of heart failure. *Heart Lung*. 1998;27:3-21.
7. Francis GS, Gassler JP, Sonnenblick EH. Pathophysiology and diagnosis of heart failure. In: Fuster V, Alexander RW, O'Rourke RA, Roberts R, King SB III, Wellens HJJ, eds. *Hurst's The Heart*. 10th ed. New York, NY: McGraw-Hill; 2001:655-685.
  8. Fonarow GC. Strategies to improve the use of evidence-based heart failure therapies: OPTIMIZE-HF. *Rev Cardiovasc Med*. 2004;5(suppl 1):S45-S54.
  9. Gammonley J, Howie A, Kirwin S, et al. *Animal-Assisted Therapy: Therapeutic Interventions*. Renton, WA: Delta Society; 1997.
  10. Allen K, Blascovich J, Tomaka J, Kelsey R. Presence of human friends and pet dogs as moderators of autonomic responses to stress in women. *J Pers Soc Psychol*. 1991;61:582-589.
  11. Grossberg JM, Alf EF Jr. Interaction with pet dogs: effects on human cardiovascular response. *J Delta Soc*. Winter 1985;2(1):20-27.
  12. Todd-Schuelke S, Trask B, Wallace C, Baun MM, Bergstrom N, McCabe B. Physiological effects of the use of a companion animal dog as a cue to relaxation in diagnosed hypertensives. <http://www.deltasociety.org/AnimalsHealthAdultsPhysiological.htm>. Accessed July 29, 2007.
  13. Wilson CC. Physiological responses of college students to a pet. *J Nerv Ment Dis*. 1987;175:606-612.
  14. Friedmann E, Katcher AH, Thomas SA, Lynch JJ, Messent PR. Social interaction and blood pressure: influence of animal companions. *J Nerv Ment Dis*. 1983;171(8):461-464.
  15. Barker SB, Dawson KS. The effects of animal-assisted therapy on anxiety ratings of hospitalized psychiatric patients. *Psychiatr Serv*. 1998;49(6):797-801.
  16. Barker SB, Pandurang AK, Best ALM. Effects of animal-assisted therapy on patients' anxiety, fear, and depression before ETC. *J ETC*. 2003;19(1):38-44.
  17. Wilson CC. The pet as an anxiolytic intervention. *J Nerv Ment Dis*. 1991;179:482-489.
  18. Cole KM, Gawlinski A. Animal-assisted therapy: the human-animal bond. *AACN Clin Issues*. 2000;11:139-149.
  19. Holcomb R, Meachman M. Effectiveness of an animal-assisted therapy program in an inpatient psychiatric unit. *Anthrozoös*. 1989;2:259-264.
  20. Raina P, Walter-Toews D, Bonnet B, Woodward C, Abernathy T. Influence of companion animals on the physical and psychological health of older people: an analysis of a one-year longitudinal study. *J Am Geriatr Soc*. 1999;47:323-329.
  21. Fila D. The significance of companion animals to a geriatric vascular patient: a case study. *Holist Nurs Pract*. 1991;5(2):11-15.
  22. Allen K, Shykoff BE, Izzo JL Jr. Pet ownership, but not ACE inhibitor therapy, blunts home blood pressure responses to mental stress. *Hypertension*. 2001;38(4):815-820.
  23. Friedmann E, Thomas SA. Pet ownership, social support, and one-year survival after acute myocardial infarction in the Cardiac Arrhythmia Suppression Trial (CAST). *Am J Cardiol*. 1995;76(17):1213-1217.
  24. Friedmann E, Katcher AH, Lynch JJ, Thomas SA. Animal companions and one-year survival of patients after discharge from a coronary care unit. *Public Health Rep*. 1980;95(4):307-312.
  25. Friedmann E, Thomas SA, Stein PK, Kleiger RE. Relation between pet ownership and heart rate variability in patients with healed myocardial infarcts. *Am J Cardiol*. 2003;91(6):718-721.
  26. Moser DK, Worster PL. Effect of psychosocial factors on physiologic outcomes in patients with heart failure. *J Cardiovasc Nurs*. 2000;14:106-115.
  27. Piano MR, Kim SD, Jarvis C. Cellular events linked to cardiac remodeling in heart failure: targets for pharmacologic intervention. *J Cardiovasc Nurs*. 2000;14:1-23.
  28. Chiariello M, Perrone-Filardi P. Pathophysiology of heart failure. *Miner Electrolyte Metab*. 1999;25(1-2):6-10.
  29. Allen K, Blascovich J, Mendes W. Cardiovascular reactivity and the presence of pets, friends, and spouses: the truth about cats and dogs. *Psychosom Med*. 2002;64:727-739.
  30. Baun MM, Bergstrom N, Langston NF, Thoma L. Physiological effects of human/companion animal bonding. *Nurs Res*. 1984;33(3):126-129.
  31. Friedmann E, Thomas SA. Health benefits of pets for families. In: Marvin B, Sussman M, eds. *Pets and the Family*. New York, NY: Haworth Press; 1985:191-203.
  32. Katcher AH. Interactions between people and their pets: form and function. In: Fogle B, ed. *Interrelationships Between People and Pets*. Springfield, IL: Charles C Thomas Publisher; 1981:41-67.
  33. Katcher AH, Friedmann E. Potential health value of pet ownership. *Continn Educ*. 1980;2:117-121.
  34. Khan MA, Farrag N. Animal-assisted activity and infection control implications in a healthcare setting. *J Hosp Infect*. 2000;46(1):4-11.
  35. Zisselman MH, Rovner BW, Schmuely Y, Ferrie P. A pet therapy intervention with geriatric psychiatry inpatients. *Am J Occup Ther*. 1996;50(1):47-51.
  36. Straatman I, Hanson EKS, Endenburg N, Mol JA. The influence of a dog on male students during a stressor. *Anthrozoös*. 1997;10(4):191-197.
  37. Grossberg JM, Alf AF Jr, Vormbrock JK. Does pet presence reduce human cardiovascular responses to stress? *Anthrozoös*. 1988;2:38-44.
  38. Schantz P. Preventing potential health hazards incidental to the use of pets in therapy. *Anthrozoös*. 1990;4:14-23.
  39. Jorgenson J. Therapeutic uses of companion animals in health care. *Image J Nurs Sch*. 1997;29(3):219-254.
  40. Lerner-DurJawa L. Pet visitation: is an infection control issue? [abstract] *Am J Infect Cont*. 1994;22:112.
  41. Cole KM, Gawlinski A. Animal-assisted therapy in the intensive care unit: a staff nurse's dream comes true. *Nurs Clin North Am*. 1995;30:529-537.
  42. Borow KM, Newburger JW. Noninvasive estimation of central aortic pressure using the oscillometric method for analyzing systemic artery pulsatile blood flow: comparative study of indirect systolic, diastolic, and mean brachial artery pressure with simultaneous direct ascending aortic pressure measurements. *Am Heart J*. 1982;103(5):879-886.
  43. Kadota L. Theory and application of thermodilution cardiac output measurement: a review. *Heart Lung*. 1985;14:605-616.
  44. Jiang NS, Machacek D, Wadel OP. Further study on the two-column plasma catecholamine assay. *Mayo Clin Proc*. 1976;51:112-116.
  45. Spielberger CD, Gorsuch RL, Lushene RE. *Test Manual for the State-Trait Anxiety Inventory*. Palo Alto, CA: Consulting Psychologists Press; 1972.
  46. Anderson WP, Reid CM, Jennings GL. Pet ownership and risk factors for cardiovascular disease. *Med J Aust*. 1992;157:298-301.
  47. Thomas SA, Liehr P. Cardiovascular reactivity during verbal communication: an emerging risk factor. *J Cardiovasc Nurs*. 1995;9:1-11.
  48. Thomas SA, Freed CD, Freidmann E, Stein R, Lynch JJ, Rosch PJ. Cardiovascular responses of patients with cardiac disease to talking and exercise stress testing. *Heart Lung*. 1992;21:64-73.
  49. Nemens E, Woods SL. Normal fluctuations in pulmonary artery and pulmonary capillary wedge pressures in acutely ill patients. *Heart Lung*. 1982;11(5):393-398.
  50. Moser DK, Dracup K, Woo MA, Stevenson LW. Voluntary control of vascular tone by using skin-temperature biofeedback-relaxation in patients with advanced heart failure. *Altern Ther Health Med*. 1997;3(1):51-59.
  51. Nicholls DP, Onuoha GN, McDowell G, et al. Neuroendocrine changes in chronic cardiac failure. *Basic Res Cardiol*. 1996;91(suppl 1):13-20.
  52. Packer M. The neuroendocrine hypothesis: a therapy to explain the mechanism of disease progression in heart failure. *J Am Cardiol*. 1992;20:248-254.
  53. Eliot RS. Relationship of emotional stress to the heart. *Heart Dis Stroke*. 1993;2(3):243-246.
  54. Lacey JI, Kagen J, Lacey BC, Moss HA. Situational determinants and behavioral correlates of autonomic response patterns. In: Knapp PH, ed. *Expression of the Emotions in Man*. New York, NY: International University Press; 1963:161-197.

---

To purchase electronic or print reprints, contact The InnoVision Group, 101 Columbia, Aliso Viejo, CA 92656. Phone, (800) 809-2273 or (949) 362-2050 (ext 532); fax, (949) 362-2049; e-mail, [reprints@aacn.org](mailto:reprints@aacn.org).

**CE Test** Test ID A0716063: **Animal-Assisted Therapy in Patients Hospitalized With Heart Failure.** *Learning objectives:* 1. Identify the physiological findings associated with advanced heart failure. 2. Discuss the overall effects of animal-assisted therapy on cardiopulmonary pressures, neurohormonal levels, and anxiety in advanced heart failure patients who participated in this study. 3. Describe the indications for further research in animal-assisted therapy that this study identifies.

**1. A stress-induced increase in epinephrine level directly facilitates release of which neurohormone?**

- a. Aldosterone
- b. Dopamine
- c. Angiotensin
- d. Norepinephrine

**2. Which of the following most contributes to progression of heart failure to advanced heart failure?**

- a. Chronic neurohormonal activation
- b. Deleterious activation of the parasympathetic nervous system
- c. Chronic hypertension
- d. Left ventricular dysfunction

**3. What is the most significant potential hazard associated with animal-assisted therapy (AAT)?**

- a. Patients' perception of the animal as a source of stress
- b. Transmission of zoonotic infections
- c. Patients' indifference to the animal
- d. An allergen-antibody response to the therapy animal

**4. What measure was implemented to ensure validity and reliability of the plasma levels of epinephrine and norepinephrine specifically?**

- a. All measuring instruments were leveled, zeroed, and calibrated prior to each use.
- b. Blood samples were collected by specially trained laboratory personnel.
- c. Intravenous infusions of drugs were not manipulated and oral medications were not given 1 hour before or after blood sampling.
- d. Patients were supine with the head of the bed elevated 45° for 15 minutes prior to and during blood sample collection.

**5. Which of the following is true regarding the data collectors used in this study?**

- a. They were specifically trained in collection of hemodynamic variables.
- b. They were non-nurse research professionals.
- c. They used specific scripting to provide explanations to each patient.
- d. They worked in pairs consisting of a research assistant and the study's coinvestigator.

**6. Advanced heart failure is indicated by which of the following?**

- a. Elevated physiological stress of living with a chronic life-threatening illness and frequent hospitalizations
- b. Accompanying pulmonary hypertension
- c. Presence of ischemic cardiomyopathy
- d. Systolic and diastolic left ventricular dysfunction

**7. Blood pressure was measured automatically by using a bedside monitor for what reason?**

- a. Manual blood pressure measurements are too time consuming.
- b. Reliability and validity of noninvasive automatic blood pressure measurements have been established.
- c. Disparities in measurements due to data collector variances are avoided when automatic monitor measurements are used.
- d. Manual blood pressure measurements themselves have been shown to increase state anxiety.

**8. Measurements of which dependent variable were completed twice rather than the 3 times accomplished for the other variables?**

- a. Cardiac index
- b. Pulmonary capillary wedge pressure
- c. State anxiety
- d. Systemic vascular resistance

**9. Which of the following findings would have excluded a patient's participation in this study?**

- a. Pre-study presence of an indwelling pulmonary artery catheter
- b. Fear of dogs
- c. White blood cell count of less than 4500 cells/μL or greater than 11 000 cells/μL
- d. Significantly advanced coronary artery disease

**10. Which baseline parameter was the only one to differ significantly between groups of patients in this study?**

- a. Dog ownership
- b. Pulmonary capillary wedge pressure
- c. Numbers and types of medications currently being taken
- d. Patients' class according to the New York Heart Association functional classification

**11. Which of the following is a hallmark of the deleterious neuroendocrine cascade that occurs in patients with advanced heart failure?**

- a. Increased myocardial cell death
- b. Loss of balance between the release of hormones from sympathetic nerve terminals and their reuptake and degradation
- c. Blunted inotropic effects of endogenous catecholamines
- d. Increased plasma levels of catecholamines

**12. Which of the following is the most accurate explanation for the improved hemodynamic and neuroendocrine measurements found in patients who received AAT?**

- a. AAT provides patients a meaningful stimulus and diverts attention away from the immediate stressors.
- b. AAT heightens arousal of the autonomic nervous system.
- c. AAT elevates cardiovascular reactivity in patients with cardiovascular disorders.
- d. AAT stimulates the release of brain neuropeptides, which in turn decrease heart rate and blood pressure.

**13. Limitations of this research study include which of the following?**

- a. Study population size and patient fluid limitations
- b. Number of facilities where study conducted and percentage of patients in the study population who were dog owners
- c. Patients' fluid limitations and percentage of patients in the study population who were dog owners
- d. Number of facilities where study conducted and amount of time patients were exposed to AAT

Test ID: A0716063 Contact hours: 1.0 Form expires: November 1, 2009. Test Answers: Mark only one box for your answer to each question. You may photocopy this form.

1. a b c d    2. a b c d    3. a b c d    4. a b c d    5. a b c d    6. a b c d    7. a b c d    8. a b c d    9. a b c d    10. a b c d    11. a b c d    12. a b c d    13. a b c d

Fee: AACN members, \$0; nonmembers, \$10 Passing score: 10 correct (77%) Category: A, Synergy CERP A Test writer: Ann Lystrup, RN, BSN, CEN, CFRN, CCRN

AMERICAN ASSOCIATION of CRITICAL-CARE NURSES

For faster processing, take this CE test online at [www.ajconline.org](http://www.ajconline.org) ("CE Articles in This Issue") or mail this entire page to: AACN, 101 Columbia, Aliso Viejo, CA 92656.

**Program evaluation**

- |  |                          |                          |
|--|--------------------------|--------------------------|
|  | Yes                      | No                       |
| Objective 1 was met  | <input type="checkbox"/> | <input type="checkbox"/> |
| Objective 2 was met  | <input type="checkbox"/> | <input type="checkbox"/> |
| Objective 3 was met  | <input type="checkbox"/> | <input type="checkbox"/> |
| Content was relevant to my nursing practice  | <input type="checkbox"/> | <input type="checkbox"/> |
| My expectations were met   | <input type="checkbox"/> | <input type="checkbox"/> |
| This method of CE is effective for this content  | <input type="checkbox"/> | <input type="checkbox"/> |
| The level of difficulty of this test was:  |                          |                          |
| <input type="checkbox"/> easy <input type="checkbox"/> medium <input type="checkbox"/> difficult |                          |                          |
| To complete this program, it took me _____ hours/minutes.  |                          |                          |

Name \_\_\_\_\_ Member # \_\_\_\_\_  
 Address \_\_\_\_\_  
 City \_\_\_\_\_ State \_\_\_\_\_ ZIP \_\_\_\_\_  
 Country \_\_\_\_\_ Phone \_\_\_\_\_ E-mail address \_\_\_\_\_  
 RN License #1 \_\_\_\_\_ State \_\_\_\_\_  
 RN License #2 \_\_\_\_\_ State \_\_\_\_\_  
 Payment by:  Visa  M/C  AMEX  Check  
 Card # \_\_\_\_\_ Expiration Date \_\_\_\_\_  
 Signature \_\_\_\_\_

The American Association of Critical-Care Nurses is accredited as a provider of continuing nursing education by the American Nurses Credentialing Center's Commission on Accreditation. AACN has been approved as a provider of continuing education in nursing by the State Boards of Nursing of Alabama (#ABNP0062), California (#01036), and Louisiana (#ABN12). AACN programming meets the standards for most other states requiring mandatory continuing education credit for relicensure.