Quantitative Evaluation of the First Order Creatine-Kinase Reaction Rate Constant in in vivo Shunted Ovine Heart Treated with Oxandrolone Using Magnetization Transfer $^{31}$P Magnetic Resonance Spectroscopy (MT-$^{31}$P-MRS) and $^{1}$H/$^{31}$P Double-Tuned Surface Coil: a Preliminary Study

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Purpose: Children born with single ventricle physiology demonstrate poor growth rate and suffer from malnutrition, which lead to increased morbidity and mortality in this population. We assume that an anabolic steroid, oxandrolone, will promote growth in these infants by improving myocardial energy utilization. The purpose of this paper is to study the efficacy of oxandrolone on myocardial energy consumption in these infants.

Materials and Methods: We modeled single ventricle physiology in a lamb by prenatally shunting the aorta to the pulmonary artery and then postnatally, we monitored cardiac energy utilization by quantitatively measuring the first order reaction rate constant, $k_f$, of the creatine-kinase reaction in the heart using magnetization transfer $^{31}$P magnetic resonance spectroscopy, home built $^{1}$H/$^{31}$P transmit/receive double tuned coil, and transmit/receive switch. We also performed cine MRI to study the structure and dynamic function of the myocardium and the left ventricular chamber. The spectroscopy data were processed using home-developed python software, while cine data were analyzed using Argus software.

Results: We quantitatively measured both the first order reaction rate constant and ejection fraction in the control, shunted, and the oxandrolone-treated lambs. Both $k_f$ and ejection fraction were found to be more significantly reduced in the shunted lambs compared to the control lambs, and they are increased in oxandrolone-treated lambs.

Conclusion: Some improvement was observed in both the first order reaction rate constant and ejection fraction for the lamb treated with oxandrolone in our preliminary study.

Keywords: Single ventricle (SV) physiology; T/R switch; High-power radiofrequency (RF) switch; First order reaction rate ($k_f$); Ejection fraction (EF); Magnetic resonance imaging (MRI); Magnetic resonance spectroscopy (MRS)
INTRODUCTION

Infants with single ventricle (SV) physiology account for a significant proportion of child healthcare resources, but tools for efficient comparison of new or even standard treatment strategies are lacking. Multiple specific types of congenital cardiac malformations result in single ventricle physiology, including hypoplastic left heart syndrome and tricuspid atresia. The vast majority of SV infants require a series of two to three palliative surgeries (Fontan) (1) because the heart cannot be surgically corrected in ways to allow for normal cardiac physiology. SV infants have a mortality rate of approximately 25% in the first year of life and survivors experience a significant lifetime morbidity rate. Children, even those with excellent technical repairs, may slowly deteriorate in the postoperative period from multiple etiologies including wound infections, prolonged effusions, prolonged mechanical ventilation and venous thrombosis. Growth failure and malnutrition are complications common to virtually all of these populations (2), and poor growth is associated with increased morbidity and mortality (3). The growth failure factor found in SV infants may be due to the high cardiac energy requirements from their shunt dependent physiology (4, 5). The growth rate has been difficult to modify, and simply increasing the caloric intake or changing the method of feeding has not proven uniformly successful (6). We hypothesize that an anabolic steroid, oxandrolone will promote growth in SV infants by improving myocardial energy utilization.

Oxandrolone is approved to promote growth in children with histories of extensive surgery, severe trauma or burns, chronic infections, constitutional growth delay and those afflicted with Turner’s syndrome (7–10). The most extensive clinical experience with the therapeutic use of oxandrolone may be seen in pediatric burn patients (11–13), in whom oxandrolone has been safely administered for one year, with beneficial effects on lean body mass and bone mineral content identifiable five years post-treatment (8). Echocardiographic evaluation in this study suggests more efficient myocardial energy usage in patients on oxandrolone, compared to the placebo group at the completion of therapy, and at one year following completion (8). Oxandrolone, utilized as a therapy to improve metabolic efficiency, has not yet been implemented in the treatment of newborn infants or patients with congenital heart defects (CHD).

To investigate the effects of oxandrolone on myocardial energy consumption in the SV infants, we developed a neonatal animal model in which SV physiology was mimicked in a lamb by prenatally shunting the aorta to the pulmonary artery (14). Sheep are the species of choice for development of this model because newborn lambs are of similar size (3–5 kg) to human newborns, pregnant ewes typically carry one to three fetuses per pregnancy and both ewes and fetuses are tolerant of the effects of surgical intervention. After birth, the lambs were treated with oxandrolone for the first month of life. The change in myocardial energy utilization with oxandrolone treatment of the lambs was determined by measuring the first order reaction rate (k_f) of the creatine-kinase (CK) reaction in the heart using magnetization transfer \(^{31}\text{P}\)-MR spectroscopy (MT-\(^{31}\text{P}\)-MRS) (15). We also performed cine MRI on the lamb to study the structure and dynamic function of the myocardium and the loading of the left ventricle by evaluating end diastolic volume (EDV), end systolic volume (ESV), stroke volume (SV) and ejection fraction (EF).

\(^{31}\text{P}\) spectroscopy allows non-invasive monitoring of phosphorous metabolites, and is being investigated as a tool to scrutinize the bioenergetics of many body organs in conditions of both health and disease (15). Several \(^{31}\text{P}\) MRS techniques, such as conventional saturation-transfer MRS (ST MRS) (16), four-angle ST (FAST) (17), three acquisition ST (novel strategy method) (18), triple repetition time ST (TRiST) (19), two repetition time ST (TwiST) (20), time-dependent ST (TDST) (21) and Bloch-Siegert four angle ST (BOAST) (22) have been employed to evaluate myocardial pseudo first order rate constant of creatine-kinase reaction in human and animal hearts.

In the present work we adopted three acquisition ST methodology in one-dimensional MT-\(^{31}\text{P}\)-CSI pulse sequence to study the myocardial energy metabolism in animal models of congenital cardiac malformations.

Theory

The CK reaction is essential for the buffering and rapid regeneration of adenosine triphosphate (ATP) in heart tissue (21). The heart uses this energy to contract. The CK enzyme catalyzes the reaction, in which the exchange of phosphate between ATP and phosphocreatine (PCr) takes place according to the reaction:

\[
\text{PCr}^+ + \text{ADP} + \text{H}^+ \rightarrow \text{Cr} + \text{ATP}^2. \tag{1}
\]

The reaction proceeds in both the forward and reverse directions with \(k_f\) and \(k_r\) as the pseudo first-order forward and reverse rate constants, respectively. One approach to
the quantification of $k_f$ is to saturate $\gamma$-ATP so that the reverse reaction brings no magnetization from $\gamma$-ATP to PCr. In MT-$^{31}$P-MRS, a train of sinc radiofrequency (RF) pulses with a narrow bandwidth (75 Hz) is applied to saturate $\gamma$-ATP line. The rate equation of longitudinal magnetization of PCr during these saturation pulses is given by modified Bloch equation:

$$\frac{dM_{PCr}(t)}{dt} = \frac{M_{ss}^{PCr} - M_{PCr}(t) - k_f M_{PCr}(t)}{T_1}. \tag{2}$$

The detailed solution of Equation 2 can be found elsewhere, in (23-25). The expression of $k_f$ obtained by solving this first order differential equation can be written as:

$$k_f = R'(1 - \frac{M_{ss}^{PCr}}{M_{ss}^{PCr}}), \tag{3}$$

where the apparent relaxation rate, $R'$, measured using $\gamma$-ATP saturated is given by the following expression:

$$R' = R_1 + k_f = 1 \ln \left(1 - \frac{M_{ss}^{PCr}}{M_{ss}^{PCr}}\right), \tag{4}$$

where, $M_{ss}^{PCr}$ is the PCr magnetization obtained with a short TR and continuous saturation pulses at $\gamma$-ATP and $R_1 = 1/T_1$ is the intrinsic relaxation rate.

Since $\gamma$-ATP is separated from PCr only by 2.5 ppm which is about 125 Hz at 3T, the RF pulse used to selectively saturate $\gamma$-ATP also partially saturates PCr peak. The rate equation of longitudinal magnetization of PCr incorporating the RF bleed-over effect in the CK reaction is given by the modified Bloch equation (26):

$$\frac{dM_{PCr}(t)}{dt} = \frac{M_{ss}^{PCr} - M_{PCr}(t) - k_f M_{PCr}(t) - \eta M_{PCr}(t)}{T_1}. \tag{5}$$

Herein, $\eta$ is the rate of loss of PCr magnetization due to an RF saturation pulse. The value of $\eta$ can be estimated by measuring the PCr magnetizations with saturation pulse irradiated at +2.5 ppm and +20 ppm on the opposite side of $\gamma$-ATP. The expression of RF bleed-over obtained by solving equation 5 can be written as:

$$\eta = \frac{R''_1 M_{ss}^{PCr}}{M_0} \left(\frac{M_{ss}^{PCr}}{M_0} - 1\right), \tag{6}$$

where $M_0^+$ and $M_0$ are the thermal equilibrium magnetization with saturation pulse irradiated at +2.5 ppm and +20 ppm, respectively and $R''_1 = R_1 + k_f + \eta$.

**MATERIALS AND METHODS**

**Animal Model**

Both timed pregnant ewe (that was preferably carried twins/triplets) and the fetuses were anesthetized at approximately 140 days of gestation (term gestation 145 days). The ewe underwent an abdominal incision, and then a uterine incision to allow exposure of the left forelimb and chest of the fetus. A left lateral thoracotomy was performed on the fetus and a 2 mm length of 8.0 mm diameter Gore-Tex vascular graft (a gift of W. L. Gore & Associates, Inc.) was interposed between the ascending aorta and main pulmonary artery. All incisions were closed, and the ewe was awakened from anesthesia. Antibiotics were given to the ewe, intraoperatively and then on a daily basis thereafter, until two days after delivery. Lambs were allowed to deliver spontaneously, and were cared for by the ewe until approximately one month of age (when MR studies were performed). The lambs rejected by the mother were bottle fed. Unoperated twins of the shunted lambs served as controls. Four groups of total nine lambs (~two lambs per group): control (n = 3), shunted (n = 2), oxandrolone-treated (ox-treated) control (n = 2), and ox-treated shunt (n = 2) were created. The ox-treated animals were given 0.1 mg/kg orally twice daily beginning on Day of Life 2. All animal work was performed under a protocol approved by the local Institutional Animal Care and Use Committee (IACUC) and euthanasia was performed in accordance with American Veterinary Medical Association (AVMA) requirements.

**Pulse Sequence**

Siemens’ product pulse sequence, CSI-free induction decay (FID) was modified to implement one-dimensional MT-$^{31}$P -CSI pulse sequence by adding outer-volume-suppression (OVS) pulses with user-selectable number of iterations, adiabatic-half-passage (AHP) RF pulse for excitation, a train of narrowband RF pulses to saturate $\gamma$-ATP, and 1D CSI with correct localization to the existing
functionalities using the pulse sequence development environment software (IDEA, Siemens Medical Solutions, Erlangen, Germany). The schematic timing diagram of MT-\(^{31}\)P-CSI is shown in Figure 1.

The 1D MT-CSI protocol for the lamb's heart is detailed in Table 1. Three hard pulses of 400 μs duration and a flip angle of 110° followed by a spoiling gradient were applied to null the longitudinal magnetizations of all metabolites. A sinc RF pulse with bandwidth 75 Hz and flip angle 300° was used as a MT saturation pulse to saturation the Y-ATP for the duration \(t_{\text{sat}}\) by repetitive application so that the reverse transfer of magnetization from Y-ATP to PCr could not proceed. Six OVS pulses and gradients were applied to prevent the contamination on the \(^{31}\)P signal of the heart from the \(^{31}\)P signal of the chest muscle. An AHP pulse of bandwidth 4 kHz was used for excitation.

### Hardware

Most clinical MR systems have a single RF transmit port for both \(^1\)H and \(^{31}\)P frequencies. A high power RF switch is positioned prior to the \(^1\)H and \(^{31}\)P TR switches as shown in Figure 2 for the automatic switching of the RF pulses to either the \(^1\)H or \(^{31}\)P TR switch for \(^1\)H MRI or \(^{31}\)P MRS, respectively.

### \(^1\)H/\(^{31}\)P Dual-Tuned RF Coils

Due to a low concentration and relatively small gyromagnetic ratio of \(^{31}\)P in human and animal tissue compared to protons (27, 28), a dedicated \(^{31}\)P coil optimized for the \(^{31}\)P nucleus accompanied by a \(^1\)H coil for the scout imaging and shimming of the region of interest is required. We built a quadrature \(^{31}\)P coil with dimensions optimized to the heart of lambs aged four to six weeks, and a relatively large \(^1\)H linear coil. The detailed explanation for the construction of the coil is given in the author's previous paper (29). The \(^{31}\)P coil consisted of a butterfly loop decoupled with a rectangular loop for a quadrature mode as shown in Figure 3, by adjusting the overlapping

| \(M_0\) | \(\Delta f_{\text{MT}}\) (ppm) | TR (s) | \(N_{\text{Sat}}\) | \(t_{\text{preSat}}\) (s) | \(t_{\text{Sat}}\) (s) |
|-----|-------|------|------|----------------|----------------|
| \(M_{\text{PCr}}\) | 0 | 20 | 0 | 0 | 0 |
| \(M_{\text{PCr}}\) | -2.7 | minimum | 0 | 2.5 | 3.5 |
| \(M_{\text{Sat}}\) | -2.7 | minimum | 2 | 0 | 2 |

CSI = chemical shift imaging; MT = magnetization transfer; TR = repetition time

Fig. 1. Timing diagram of pulse sequence, 1D MT-CSI.
region of the coils. The $^1$H RF coil that surrounded the $^{31}$P coil, as shown in the Figure 3, was positioned such that the field produced by the coil would be perpendicular to that produced by the $^{31}$P butterfly coil. The coupling between the $^1$H and $^{31}$P rectangular coil was minimized by choosing the large dimension of the $^1$H coil compared to the rectangular coil. The loaded isolation between the $^1$H and $^{31}$P butterfly coils measured using vector network analyzer (E5061B ENA Series, Agilent Technologies, Englewood, CO, USA) and 250 mL saline bag phantom at the $^1$H frequency was -23.2 dB and that at the $^{31}$P frequency was -28.7 dB while that between the $^1$H and $^{31}$P rectangular coils at the $^1$H frequency was -22.8 dB and at the $^{31}$P frequency was -26.7 dB. Moreover, the $^{31}$P coil affects the $^1$H signal to noise ratio (SNR) far more than the $^1$H coil affects the $^{31}$P SNR (30). This is because $^{31}$P coils have relatively high capacitor values compared to the $^1$H coil, which provides low impedance at the $^1$H frequency (31). The isolation between the $^1$H and $^{31}$P RF coils at $^1$H frequency could be further increased by inserting the $^1$H traps at the $^{31}$P coils with a little compromise of the $^{31}$P signal. No extra lossy components on the $^{31}$P coil were added so that the performance of the $^{31}$P coil is same as the same coil on the same frame without $^1$H coil. The separate coil design for each nucleus allows us to

Fig. 2. Block diagram of hardware.

Fig. 3. (a) Circuit diagram and (b) photograph of $^1$H/$^{31}$P dual tuned RF coils. The $^{31}$P loop consisted of a butterfly loop and an inner rectangular loop decoupled geometrically for a quadrature mode. The outer rectangular loop was $^1$H RF coil whose surface area was made large to minimize the coupling with $^{31}$P rectangular loop, and positioned such that it was in quadrature mode with $^{31}$P butterfly loop.
independently match and tune the $^1$H and $^{31}$P coils without interference.

**T/R and High Power RF Switches**

The T/R circuitry for our $^1$H/$^{31}$P dual-tune RF coil consists of a linear T/R switch for the $^1$H coil and a quadrature T/R switch for the $^{31}$P coil with high-power passive RF switch (29, 32) consisting of two parallel LC trap circuits placed prior to the T/R switches to automatically direct the $^1$H RF pulse to the $^1$H T/R switch during $^1$H MRI and the $^{31}$P RF pulse to the $^{31}$P T/R switch during $^{31}$P MRS.

**MRI Experiments**

The MT-$^{31}$P MRS experiments were carried out on nine lambs aged four to six weeks and in four groups: control ($n = 3$), ox-treated control ($n = 2$), shunted ($n = 2$) and ox-treated shunted ($n = 2$), using the custom $^{31}$P RF pulse to the $^1$H T/R switch during $^1$H MRI and the $^{31}$P RF pulse to the $^{31}$P T/R switch during $^{31}$P MRS.

1H MR anatomical images were first acquired with the lamb in the scanner (in a feet first lateral position) to determine the position of RF coils using the localizer. A $^1$H marker attached to the center of the $^1$H/$^{31}$P coil was used to adjust coil at the position of heart. The ovine heart was then shimmmed to homogenize the $B_0$ field over the region of heart. Cardiac-triggered, 1D CSI with eight phase encodes and saturation bands on the chest to minimize contamination from the chest muscle were used for $^{31}$P spectroscopy. Three experiments were carried out to measure $k_f$. $M_{po}^{o}$ was obtained using TR = 20 sec with no MT and $M_o$, saturation pulses, $M_{po}^{TR}$ using three hard RF pulses with flip angle 300° as $M_o$, saturation RF and a spoiling gradient pulses followed by MT saturation pulses at -2.7 ppm away from PCr (at the position of $\gamma$-ATP) for 2 sec, and $M_{pc}^{o}$, was measured with pre-saturation delay of 2.1 sec and MT saturation at -2.7 ppm for 3.5 second (18). The total scan time for measuring $M_{po}^{TR}$, $M_o^{TR}$, and $M_{pc}^{TR}$ was 37 minutes 40 seconds. The value of RF bleed over was obtained by measuring $M_o$ and $M_o^{TR}$ and using equation [6]. The thermal equilibrium magnetization, $M_o^{o}$ under RF bleed-over was measured by irradiating saturation RF pulses at +2.5 ppm for $t_{sat}$ = 20 s. The same experiment was repeated to measure thermal equilibrium magnetization, $M_o$ without RF bleed-over effect by irradiating saturation RF pulses at +20 ppm (26). All experiments were performed with a receiver bandwidth of 2.5 kHz and 512 data points.

Only four out of the nine lambs (i.e. one in each group of control, shunted, ox-treated control, and ox-treated shunted lambs) underwent cine MRI after $^{31}$P MRS using the standard Siemens’ coil (Flex Large 4, a 3T Tim) as a receiver and the body coil as a transmitter. Long-axis and short axis scout images were acquired for planning short axis cine images. Cine images covering the whole heart in short axis orientation were acquired using TRUIFI (true fast imaging with steady state free precession) pulse sequence with retrospective ECG gating. The imaging parameters were: typical field of view (FOV) = 260 × 142 mm², typical matrix size 208 × 114, in-plane resolution = 1.25 × 1.25 mm, contiguous stack of slices of 6 mm thickness, TE/TR = 1.48 ms/3.4 ms, flip angle = 40°, temporal resolution = 23.6 ms.

The raw data for the $^{31}$P MRS was processed using home-developed processing software in Python language that performs 10 Hz Gaussian and Lorentzian broadening, zero filling to 4-k of data points, fast Fourier transformation, and interactive zeroth and first order phase correction. The magnetization of PCr and ATP in myocardium was measured as the area under the respective resonance peaks.

The cine data were processed manually using Argus software (Siemens Healthcare) to calculate end diastolic volume (EDV), end systolic volume (ESV), stroke volume (SV) and ejection fraction (EF). The papillary musculature was excluded from the calculation.

**RESULTS**

Figure 4a shows an example of localization image with 1D CSI grid (white) and green rectangle as shim region of in vivo lamb heart obtained with the custom made $^1$H/$^{31}$P double tuned RF coils. Figure 4b-e are the representative stacked plots of $^{31}$P MR spectra from control (triplet), shunted (triplet), ox-treated shunted (doublet) and the ox-treated control (doublet) lambs, respectively. Blue, green and red spectra indicate $M_{po}^{o}$, $M_o^{o}$, and $M_{pc}^{o}$, respectively. The values of $k_f$, SNR of the $M_o^{o}$ peak obtained from these spectra, EDV, ESV, SV, and EF are listed in Table 2. The low value of SNR of $M_{pc}^{o}$ reflects the poor shimming compared to the higher value. The $k_f$ of the triplet shunted lamb is
Fig. 4. (a) A scout image with 1D CSI grid with green rectangle as a shim region. Saturation bands were used on the chest wall to avoid the contamination on the $^{31}$P signal of heart from the chest wall. (b-e) in-vivo stacked plots of $^{31}$P MR spectra of control (triplet), shunted (twin), and ox-treated control (twin) lamb. Blue, green and red spectra indicate $M_{\text{PCr}}^0$ (equilibrium), $M_{\text{PCr}}$ (saturation pulse for 3.5 second and pre-saturation time of 2.5 second), and $M_{\text{PCr}}^\alpha$ (saturation pulse for two seconds), respectively. The arrows indicate the positions where saturation pulse is applied. (f) $^{31}$P spectra with (green) and without (blue) RF bleed-over effect. There is a decrease in PCr peak due to RF bleed-over effect.
0.09 s\(^{-1}\) which is much smaller than either of the control lambs (0.22 s\(^{-1}\) or 0.28 s\(^{-1}\)) of the same set of triplets. The \(k_f\) of the ox-treated shunted lamb of the last doublet is 0.30 s\(^{-1}\) which seems to demonstrate the increase of \(k_f\) with oxandrolone treatment toward a normal value, or its ox-treated control doublet (\(k_f = 0.31\) s\(^{-1}\)). Figure 4f is the spectra for computing the value of RF bleed-over where blue and green spectra are with and without RF bleed-over effect, respectively. The RF bleed-over was found to be 2.8%.

Table 2. \(k_f\), Signal to Noise Ratio (SNR) of \(M_{\text{PCR}}\)\(^{\text{PO}}\), End Diastolic Volume (EDV), End Systolic Volume (ESV), Systolic Volume (SV), and Ejection Fraction (EF) of Different Groups of Lambs

| Lamb type     | Lamb       | \(k_f\) (s\(^{-1}\)) | SNR (\(M_{\text{PCR}}\)\(^{\text{PO}}\)) | EDV (mL) | ESV (mL) | SV (mL) | EF (%) |
|---------------|------------|---------------------|--------------------------------------|----------|----------|---------|--------|
| Triplet (~4 weeks) | Control    | 0.22                | 49.75                                | 31.0     | 16.4     | 14.6    | 47.2   |
|               | Shunted    | 0.09                | 25.0                                 |          |          |         |        |
|               | Control    | 0.28                | 23.75                                |          |          |         |        |
| Twin (~8 weeks) | Control    | 0.36                | 71.0                                 |          |          |         |        |
|               | Shunted    | 0.28                | 67.0                                 | 103.2    | 69.0     | 34.2    | 31.3   |
| Single (~4 weeks) | Ox-treated shunt | 0.32                | 37.7                                 |          |          |         |        |
| Twin (~6 weeks) | Ox-treated control | 0.15                | 34.7                                 |          |          |         |        |
|               | Shunted stillborn | -                   |                                      |          |          |         |        |
| Twin (~5 weeks) | Ox-treated control | 0.31                | 56.6                                 | 48.6     | 25.2     | 23.4    | 48.1   |
|               | Ox-treated shunt | 0.30                | 30.5                                 | 100.5    | 62.5     | 38.0    | 37.8   |

Fig. 5. Representative short axis images of shunted lamb with (a) diastolic and (b) systolic contours (red) drawn using Argus software.
of a shunted lamb heart with red contours placed on endocardial myocardial borders from base to apex on all slices pertaining to (a) end diastole and (b) end systole. The papillary muscles are excluded from the contours. Similar contours were drawn for other lambs (not shown) in order to calculate left ventricular parameters. The EF for controlled, shunted, ox-treated control, and ox-treated shunt were 47.2%, 31.3%, 48.1%, and 37.8% respectively. The EDV of the shunted (103.2 mL) or ox-treated shunted lamb (100.5 mL) was two to three times larger than the EDV of either the control (31 mL) or the ox-treated control lamb (48.6 mL), which was expected for volume overloaded hearts.

DISCUSSION AND CONCLUSION

The RF coil and T/R switches were developed to optimize the SNR of the $^{31}$P signal and enable $^1$H/$^{31}$P dual nuclear MRI/S automatically without the need to physically move and replace either the RF coil or T/R switch during the experiment. We implemented MT saturation technique in 1D CSI pulse sequence to measure $k_f$ in different groups of ovine hearts (as mentioned above) at 3T. An AHP pulse was used to generate a uniform excitation over the region of interest using the quadrature $^{31}$P surface coil.

This study is the first to measure $k_f$ and cardiac volume in surgically created SV and ox-treated lambs. We studied oxandrolone's effect on myocardial energetics and function not because it's a new drug, but because we are proposing a novel and therapeutic use for the substance for the purpose of treating growth failure in SV infants. The $k_f$ of the in vivo ovine heart was measured using novel strategy method (three acquisition method) (18) with drastically shorter total data acquisition time compared to the conventional approach which requires progressive saturation experiments with long pre and MT saturation times to evaluate $M_{PCr}^{ss}$ and $R_0$. The PCr peak in blue spectra in Figure 4b–e, for measuring $M_{PCr}^{ss}$, was reduced in green spectra for measuring $M_{PCr}^{ss}$ due to the application of MT pulses at the position of γ-ATP peak (to suppress reverse magnetization from γ-ATP to PCr) for 3.5 seconds, and even further reduced in red spectra for measuring $M_{PCr}^{ss}$ due to short TR and MT saturation pulses (applied for two seconds), as used in this acquisition.

It is fairly reasonable to compare the $k_f$ values between the lambs of same biologic twin or triplet. The values of $k_f$ obtained from our results show that, compared with the control heart of the same twins/triplets, the forward reaction rate constant of CK reaction is significantly lower in a shunted heart, and it is increased in the ox-treated heart compared to the shunted heart which may indicate the increase in the rate of energy production in the ox-treated lamb. This result is consistent with the previous studies on in vivo human hearts, which reported substantial decreases in $k_f$ in failing hypertrophic myocardium compared to healthy myocardium (33–35) and on seven month old isolated cardiomyopathic hamster heart which showed an increase in the CK flux with enalapril-treatment (36). The value of $k_f$ for the shunted or control heart of the triplets is smaller than that in the twins. The value of $k_f$ for the control and ox-treated control lamb of the last twins of Table 2 is approximately same as expected. The $k_f$ values of shunted and control lambs of one twin or triplet is different from another twin or triplet lambs. This may be due to several reasons. The triplet shunt may not get enough nutrients than the twin shunt before and after its birth. Those lambs which are rejected by the ewe after birth were bottle-fed. One lamb may be much sicker than the other. The triplet shunt was scanned when it was ~ one month old and the twin shunt was scanned when it was ~ two months old. Chesky et al. (37) from the $^{31}$P MRS on the male Fischer rat have shown that the myocardial creatine phosphokinase activity rises from one month of age to a maximum two months of age. There may not be the same experimental situation for the lambs though we tried hard to equalize the circumstance as much as possible. It might be that they were simply scanned on different dates.

Several methods, such as cardiovascular ultrasound (38), angiocardiography (39), and computed tomography (40) have been used to quantitatively evaluate the ESV, EDV, SV, and EF of the left ventricle. These methods have certain constraints and drawbacks. For instance, computed tomography is carcinogenic to the patient, cardiovascular ultrasound method requires geometric premise, etc. As an alternative to these methods, cardiac MRI is a non-invasive method that has been frequently used as a standard tool for the cardiac volume measurements (41, 42). In the present study, we used cine MRI for the volume measurements of in vivo ovine heart. The cine MRI measurement of the ovine heart shows that EF of control lamb is greater than that of the shunted heart and is similar to that of the ox-treated control lamb. There is some improvement of EF with oxandrolone treatment, as expected. The EDV and ESV of ox-treated shunts are greater than those of the ox-treated controls, which indicates the greater remodeling in
the ventricle of the shunted lamb. Further studies on more lambs, with and without, surgically created SV physiology are needed for proper statistical analyses to confirm these findings.

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