Electricity Remodeling of the Atrium in an Anatomic Model of Atrial Flutter

Relationship Between Substrate and Triggers for Conversion to Atrial Fibrillation

Joseph B. Morton, MBBS; Melissa J. Byrne, BSc; John M. Power, BVSc, PhD; Jai Raman, MBBS, M Med; Jonathan M. Kalman, MBBS, PhD

Background—Atrial flutter (AFL) and atrial fibrillation (AF) frequently coexist, yet the specific relationship between these arrhythmias, and particularly whether sustained AFL leads to AF, is unknown.

Methods and Results—We investigated the electrophysiological consequences of chronic AFL using an ovine anatomic right atrial Y-lesion model. AFL was induced in 7 animals, and 4 remained in sinus rhythm (controls). Sheep were monitored for spontaneous conversion of AFL to AF. Six of 7 sheep sustained AFL for 28 days. In 1 of 7 sheep, spontaneous conversion of AFL to AF occurred on day 5. AFL produced a highly significant fall in right and left atrial refractoriness (AERP, \( P < 0.001 \)), with 74±10% of the reduction occurring by day 3. Right atrial conduction velocity also fell significantly (baseline 89±9 cm/s versus day 28 64±14 cm/s, \( P < 0.001 \)) but over a slower time course. AERP and conduction velocity changes coincided with a characteristic biphasic decrease and increase in the AFL cycle length. The excitable gap (percent of AFL cycle length) increased from 13±3% at baseline to 46±8% by day 28 (\( P < 0.001 \)). Sustained AF (\( > 30 \) seconds) was not inducible at baseline but after 28 days of AFL could be induced in 6 of 6 sheep by critically timed single or multiple extrastimuli delivered either in sinus rhythm or AFL. There was no significant change in any parameter in control sheep.

Conclusions—In this model, AFL produced electrical remodeling and the substrate for sustained AF. However, spontaneous conversion to AF was uncommon, and the development of AF was dependent on specific triggers. (Circulation. 2002;105:258-264.)

Key Words: atrial flutter ■ fibrillation ■ remodeling

Atrial flutter (AFL) and atrial fibrillation (AF) frequently coexist,\(^1\)\(^-\)\(^3\) although the specific relationship between these 2 arrhythmias, particularly whether sustained AFL leads to AF, is unknown. In the canine sterile pericarditis model, Ortiz and coworkers\(^4\) demonstrated that conversion of AFL to AF was associated with a decrease in the length of a functional line of block in the right atrial (RA) free wall. Less is known about the mechanism of conversion from AFL to AF in an anatomic model of AFL in which lines of block are fixed. This has potential clinical relevance to “incisional” or “scar-mediated” AFL occurring in patients after surgical repair of congenital heart disease, and perhaps in a subset of patients with typical human flutter.

The concept of atrial electrical remodeling (ER) was first advanced by Wijffels et al\(^8\) when they demonstrated that AF-induced shortening of refractoriness in goats led to perpetuation of AF (“AF begets AF”). It has also been demonstrated recently in humans that atrial ER also occurs as a result of sustained AFL.\(^6\)\(^,\)\(^7\) This has led to the hypothesis that AFL may potentially beget the substrate for AF as a result of ER.\(^7\) Whether development of substrate alone is a sufficient precondition for AFL to spontaneously convert to fibrillation is unclear. Recent studies have demonstrated the importance of pulmonary vein initiating triggers not only for the onset of paroxysmal AF\(^8\) but possibly for the conversion of flutter to fibrillation.\(^9\)

In the present study, we investigated the hypothesis that in an anatomic model of AFL, progression to sustained AF would depend on the interaction between development of suitable atrial substrate and the presence of focal triggers to initiate the arrhythmia.
Methods

Ovine Model

The study was approved by the Animal Ethics Committee of the Austin and Repatriation Medical Center and conducted in accordance with guidelines outlined in the “Position of the American Heart Association on Research Animal Use” adopted on November 11, 1984, by the American Heart Association. A surgical Y lesion was created in the RA of 11 adult sheep (mean weight 45 kg) on the basis of the technique previously described by Frame and coworkers in canines.10 Surgery was performed under general anesthesia (intravenous propofol, 2 mg/kg; ventilation with 2% halothane and oxygen) without the need for cardiopulmonary bypass.

Access to the RA and left atrium (LA) was by sequential right and left lateral thoracotomy incision. The pericardium was incised and reflected. Two full-thickness RA incisions were made from (1) inferior to superior vena cava and (2) tip of the RA appendage (RAA) to the caudal region of the intercaval incision. The connecting incisions were each closed by continuous suture to create a Y lesion.

Quadrupolar electrode plaques with adjacent pace/sense bipoles (electrode diameter 1.5 mm; 2.0 mm interelectrode distance) were sewn onto the RAA and LA appendage (LAA). A hexapolar plaque with 6 linearly arranged electrodes (electrode diameter 1.5 mm; 2.0 mm interelectrode distance) was sewn onto the RA free wall adjacent to the tricuspid annulus (TA). Wires from the electrode plaques were tunneled subcutaneously to the dorsum of the sheep, exteriorized, and attached to electrical connectors housed permanently on the animal’s back. All electrophysiological studies (EPSs) were performed via these exteriorized leads in conscious sheep.

The pericardium and chest wall were closed in layers. Intravenous antibiotics (1 g of ampicillin/80 mg of gentamicin sulfate) and an analgesic/anti-inflammatory agent (50 mg of flunixin meglumine) were administered intraoperatively and then daily for 72 hours. Animals recovered for 7 to 14 days before the experimental protocol was begun.

Protocol

Baseline Electrophysiological Evaluation

The study protocol was initiated with all animals undergoing baseline EPS.

AFL Induction

Seven animals in group 1 underwent induction of sustained AFL. Four animals in group 2 (controls) remained in sinus rhythm (SR) without AFL induction after Y-lesion creation.

Follow-Up Electrophysiological Evaluation

After initiation of the protocol, all animals underwent repeat EPS on days 3, 14, and 28. On these days, AFL was terminated by overdrive atrial pacing to allow testing during a brief period of SR. After EPS, flutter was reinitiated.

AFL Induction

AFL was induced by burst atrial pacing at a cycle length (CL) of 150 to 250 ms from the LAA electrode. AFL was defined as a macroreentrant atrial tachycardia displaying (1) typical “sawtooth” flutter wave appearance on the surface ECG, (2) entrainment criteria,11,12 with demonstration of an excitable gap, and (3) a regular and constant epicardial atrial activation pattern (ie, no change in sequence) recorded from the 3 epicardial electrode plaques.

Animals were monitored continuously with a custom-built system to document either termination of AFL or degeneration to AF. When AFL terminated, it was immediately reinitiated with programmed stimulation. Continuous atrial recordings were processed by an isolated interface/amplifier controlled by virtual instrumentation (LabView software, National Instruments) and a Macintosh computer. Every 30 seconds, the mean atrial CL was calculated during a 1-second window, and if it was >250 ms, SR was assumed and programmed atrial extrastimuli (PES) were automatically delivered to reinduce AFL. All episodes of AFL termination/reinduction were reviewed offline on a computer workstation to exclude AF and confirm AFL reinduction.

Electrophysiological Testing

A Quinton EP laboratory system (Quinton Electrophysiology Corp) was used with a programmable stimulator (Medtronic, model 5326). All measurements were made offline with screen calipers at 400 mm/s sweep speed.

Electrophysiological Parameters Measured During SR

Atrial effective refractory periods (AERPs) were performed at 4 CLs (550, 450, 350, and 250 ms) from 3 sites (RAA, LAA, and proximal bipolar of TA strip electrode). Pacing was performed at twice diastolic threshold with an 8-beat drive train and a single extrastimulus that was increased by 2-ms intervals from 70 ms. We determined the dispersion of refractoriness (AERP DISP) across the LAA/RAA/TA for each CL by subtracting the minimum from the maximum AERP.

RA conduction velocity (CV; cm/s) was determined during constant unipolar pacing at a CL of 350 ms from the proximal pole of the TA strip electrode plaque, measuring the time interval from the pacing spike to the rapid deflection of the unipolar electrogram recorded on the distal electrode. The wavelength (λ) of this impulse was also measured (λ = CV × AERP 350). RAA to LAA conduction time (ms) during constant RAA pacing at 350 ms CL was measured from RAA stimuli artifact to first recorded bipolar signal on the LAA plaque. P-wave duration (ms) was measured from lead II of the surface ECG.

Electrophysiological Parameters Measured During AFL

AFL CL was measured during AFL. In addition, excitable gap (EG) was measured. An atrial extrastimulus (S1) was delivered from the TA electrode at a coupling interval (CI) equal to the AFL CL and progressively reduced by 2-ms decrements until local refractoriness was encountered. The return cycle was measured. The EG was defined as the AFL CL minus the shortest CI that produced atrial capture, which represented the total window of reset. Resetting was
TABLE 1. Changes in AERP by pacing site and CL

<table>
<thead>
<tr>
<th>Pacing Site/CL, ms</th>
<th>AERP Baseline (n=7), ms</th>
<th>AERP Day 3* (n=7), ms</th>
<th>AERP Day 14† (n=6), ms</th>
<th>AERP Day 28 (n=6), ms</th>
<th>P (ANOVA)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LAA/550</td>
<td>132±15</td>
<td>97±21</td>
<td>82±12</td>
<td>78±16†</td>
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</tr>
<tr>
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<td>99±19</td>
<td>...</td>
<td>79±17†</td>
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<tr>
<td>LAA/350</td>
<td>143±15</td>
<td>105±19</td>
<td>...</td>
<td>80±17§</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>LAA/250</td>
<td>143±15</td>
<td>108±19</td>
<td>...</td>
<td>84±18§</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>RAA/550</td>
<td>181±23</td>
<td>108±20</td>
<td>97±18</td>
<td>96±19†</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>RAA/450</td>
<td>186±20</td>
<td>112±18</td>
<td>...</td>
<td>102±18§</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>RAA/350</td>
<td>183±20</td>
<td>119±21</td>
<td>...</td>
<td>107±19†</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>RAA/250</td>
<td>181±13</td>
<td>125±19</td>
<td>...</td>
<td>113±15§</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TA/550</td>
<td>180±19</td>
<td>114±30</td>
<td>97±15</td>
<td>92±19†</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TA/450</td>
<td>187±18</td>
<td>118±28</td>
<td>...</td>
<td>90±19†</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TA/350</td>
<td>191±16</td>
<td>121±32</td>
<td>...</td>
<td>100±17†</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TA/250</td>
<td>185±14</td>
<td>125±30</td>
<td>...</td>
<td>107±15§</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Statistics

All data are presented as mean±SD. Data measured at multiple time points were analyzed by a single-factor ANOVA with repeated measures, followed by the Tukey-Kramer procedure for multiple comparisons. Differences between 2 groups were analyzed with either Student’s paired or unpaired t test, the nonparametric Wilcoxon rank sum test for nonnormally distributed data, or χ² analysis for paired proportions. A P value <0.05 was considered statistically significant.

Results

AFL Induction

Chronic AFL was achieved in the 7 study sheep (Figures 1 and 2). In 6 of 7 sheep, AFL was sustained for 28 days without any spontaneous conversions to AF. In 1 of 7 sheep, AFL spontaneously converted to AF after 5 days of sustained AFL via a progressive shortening of the AFL CL. AF then persisted for 48 hours without reorganizing to AFL. This sheep was excluded from further analysis.

Electrophysiological Parameters Measured During SR

A significant fall in AERP occurred during 28 days of AFL (Table 1 and Figure 3a), with 74±10% of this reduction occurring by day 3. Between day 3 and day 28, there was a relative plateau in the AERP. A similar pattern was also observed for the AERP_DISP, with a statistically significant early fall and then plateau (Table 2). In the control group, there was no significant change in either AERP or AERP_DISP over the study period. The expected reduction in AERP with shortening of the drive CL was not observed at either baseline or on day 28 in either group. For all sites, mean AERPs at a 250- and 550-ms drive CL were not significantly different.
RA CV fell significantly during 28 days of AFL (baseline 89±9 cm/s versus day 28 64±12 cm/s, P<0.01) but over a slower time course than the AERP change (Figure 3b). The impulse wavelength also significantly shortened (baseline 16.9±2.0 cm, day 3 10.0±0.7 cm, day 28 6.3±1.4 cm; P<0.001). No significant change in control group CV occurred over the study period (baseline 98±6 cm/s versus day 28 105±9 cm/s, P=0.27), and the baseline difference between cases and controls was not statistically significant (P=0.14).

AFL resulted in significant increases in RAA to LAA conduction time (baseline 84±9 ms versus day 28 107±14 ms, P=0.01) and P-wave duration (baseline 48±3 ms versus day 28 66±5 ms, P=0.01). In the control group, no significant changes were observed in either RAA to LAA conduction time (baseline 86±12 ms versus day 28 85±13 ms, P=NS) or P-wave duration (baseline 47±3 ms versus day 28 48±3 ms, P=NS).

**Electrophysiological Parameters Measured During AFL**

A characteristic biphasic change in AFL CL was observed during the study period (Figure 3c; baseline 194±13 ms, day 3 160±11 ms, day 14 178±9 ms, day 28 183±8 ms; P<0.001). The early period of AFL CL shortening coincided with a fall in AERP and a 50% increase in mean EG (from 13±3% to 20±9%). The subsequent period of lengthening in the AFL CL coincided with the observation of CV slowing and AERP plateau.

The EG as measured from the TA electrode demonstrated a progressive widening (Figure 4, left) during the study period (baseline 13±3%, day 3 20±9%, day 14 38±9%, day 28 46±8%; P<0.001). By entrainment criteria (postpacing interval minus AFL CL ≤20 ms), the TA was within the AFL macroreentrant circuit, and resetting could be demonstrated in all sheep. Resetting curves constructed during the determination of the EG on day 28 showed a mixed pattern in all 6 sheep (Figure 4, right), consistent with a proportion of the EG being fully excitable (initial flat component) and a proportion being partially excitable (later increasing component).

**Spontaneous Termination of AFL**

AFL terminated spontaneously on 41 occasions in 7 sheep during the first 3 days after AFL induction, occurring predominantly (76%) over the first 24 hours (31 episodes in 7 sheep) and decreasing in frequency over day 2 (8 episodes in 4 sheep) and day 3 (2 episodes in 2 sheep). When AFL terminated spontaneously, it was immediately reinduced. After day 3, there were no further spontaneous terminations of AFL (Figure 5).

**TABLE 2. Spatial Dispersion of Refractoriness**

<table>
<thead>
<tr>
<th>Pacing CL, ms</th>
<th>Baseline (n=7), ms</th>
<th>Day 3 (n=7), ms</th>
<th>Day 28 (n=6), ms</th>
<th>P (ANOVA)</th>
</tr>
</thead>
<tbody>
<tr>
<td>550</td>
<td>57±22</td>
<td>22±16*</td>
<td>22±9†</td>
<td>0.0002</td>
</tr>
<tr>
<td>450</td>
<td>60±19</td>
<td>23±14*</td>
<td>26±9†</td>
<td>0.002</td>
</tr>
<tr>
<td>350</td>
<td>54±17</td>
<td>21±18*</td>
<td>28±15†</td>
<td>0.004</td>
</tr>
<tr>
<td>250</td>
<td>48±14</td>
<td>22±16†</td>
<td>30±7‡</td>
<td>0.016</td>
</tr>
</tbody>
</table>

*P<0.01 and †P<0.05 for comparison with baseline. ‡P=NS for comparison with day 3.

**Figure 3.** Changes in AERP, CV, and AFL CL over 28 days of AFL. a, Representative significant changes in AERP recorded from RAA, LAA, and TA electrodes at drive CL of 550 ms (P<0.001 for each site by ANOVA). There was a highly significant early fall in AERP (*) with plateau thereafter †. b, Changes in mean CV vs AFL duration for 6 sheep that sustained 28 days of AFL. Highly significant fall in CV occurred over 28 days of flutter (P=0.001 by ANOVA) but was more gradual than change in AERP (onset vs day 3, P=NS; onset vs day 28, P<0.01); c, Characteristic biphasic change in AFL CL observed over 28 days for each of 7 flutter study sheep (P<0.001 by ANOVA).
AF Vulnerability
At baseline, neither sustained AF nor sustained AFL was inducible in any sheep by a single tight-coupled extrastimulus (AF vulnerability=0%). On day 28 in the control group, sustained AF was inducible in 1 of 4 sheep (longest episode 30 seconds, AF vulnerability=2%, P=NS versus baseline), but sustained AFL was not inducible in any animal with a single extrastimulus. On day 28 in the AFL group (n=6), sustained AF was inducible by a single extrastimulus during AERP testing in each animal (AF vulnerability=40%, P<0.001 versus baseline, P<0.001 versus controls). In addition, after 28 days of AFL, a single extrastimulus delivered during AERP testing could now induce sustained AFL in 3 of 6 sheep.

In all 6 AFL sheep in which AF was induced, the arrhythmia was sustained. In 1 of these 6 sheep, AF organized to stable AFL after 24 hours, and in the remaining 5 sheep, AF persisted for 4.8±2.6 days (range 2 to 8 days) before the animals were killed, without reversion to SR or AFL.

Effect of PES on AFL
At baseline (1 hour after initial AFL induction), AFL could not be converted to sustained AF in any sheep. However, AFL was easily terminated by ≤2 PES in 7 of 7 sheep from both the RAA (1.7±0.5 extrastimuli, mean CI 164±11 ms) and LAA (1.1±0.4 extrastimuli, mean CI 140±10 ms).

On day 28, AFL could not be terminated by a single tight-coupled extrastimulus in any sheep. However, with additional PES, the following results were observed. In the RAA, 4.2±1.2 extrastimuli (P=0.03 versus baseline) delivered at a mean CI of 97±13 ms resulted in termination of AFL in 89% of episodes and conversion to AF in 11%. In the LAA, 2.8±0.7 extrastimuli (P=0.03 versus baseline) delivered at a mean CI of 87±14 ms resulted in termination of AFL in 17% of episodes and conversion to AF in 83%.

Echocardiography
There was no significant difference in LA area between the 2 groups at baseline (controls 28±4 cm², AFL sheep 26±4 cm², P=NS). By day 28, there was a significant increase in the LA area in the AFL sheep (51±6 cm², P<0.001 compared with baseline) but no change in the control group (30±2 cm², P=NS compared with baseline).

Discussion
This ovine study of AFL with a Y-lesion anatomic barrier model prospectively demonstrates that chronic AFL leads to atrial ER, with quantitative changes observed in AERP and CV. Principally by way of these changes, AFL induced the substrate for development of sustained AF not present at baseline. However, despite the development of a very short atrial wavelength, spontaneous conversion of AFL to AF occurred infrequently, and the development of AF depended on appropriate triggers (PES) either interacting with the AFL circuit or being delivered as a premature atrial extrastimulus early in the postreversion period.

Furthermore, despite the fact that it induced the substrate for AF, the stability of AFL in this anatomic model was also enhanced by the process of ER via widening of the EG. The combination of an early fall in AERP and delayed slowing of CV produced a characteristic biphasic change in AFL CL, with a consequent progressive widening of the EG. After 28 days of flutter in 6 sheep, nearly half (46±8%) of the AFL CL was excitable. Evidence for the stability of AFL after ER in this anatomic model is provided by the following: (1) All spontaneous terminations of AFL occurred within the first 3...
days after flutter initiation. (2) Multiple tightly coupled atrial extrastimuli (range 2 to 5) were required to terminate chronic AFL or to convert chronic AFL to AF. At baseline, AFL was always terminated with ≤2 extrastimuli. (3) Spontaneous conversion to AF occurred in only 1 sheep (on day 5). Thus, it might be said that in this anatomic model, "AFL begets AFL" in the absence of appropriate triggers that either lead to AF or result in AFL termination.

**AFL Model**

The relationship between AFL and AF in a functional model of AFL has been investigated extensively. In the canine sterile pericarditis model, Waldo and colleagues\(^6\) demonstrated that when lines of block are functional, stable AFL becomes a more rapid atypical flutter and then degenerates to AF when the line of block shortens. We elected to use the anatomic model of AFL described by Frame and coworkers\(^10,14,15\) to evaluate the effects of AFL on atrial remodeling and the relationship between AFL and AF when the lines of block are fixed. In that model, reentry occurs around the TA, analogous to human AFL,\(^16\) with the Y lesion providing a fixed posterior anatomic barrier. The clinical counterpart of AFL with fixed anatomic lines of block includes those patients with surgically corrected congenital heart disease (eg, atrial septal defect repair, Mustard repair, and Fontan palliation).\(^17\) In these patients, there is a high late incidence of persistent chronic AFL (or scar-mediated atrial reentry), which often remains stable for years without degeneration to AF. In typical human flutter, evidence suggests that a proportion of patients may also have fixed block along the crista terminalis,\(^18\) although in the majority, this appears to be functional\(^19\) or anisotropic. However, given the correct milieu, even a partial line of fixed block with functional extension will result in stable flutter.\(^20\)

**Atrial Remodeling and Focal Triggers**

The process of atrial arrhythmias or sustained high-rate atrial pacing that induces atrial ER has been demonstrated in numerous studies.\(^5,21–23\) A fall in effective refractory period, increased effective refractory period heterogeneity,\(^24\) and regional slowing of conduction\(^25\) have been demonstrated in response to even brief periods of AF. Fewer data exist evaluating the effects of AFL on atrial remodeling. Franz et al\(^9\) described quantitatively similar decreases in the monophasic action potential at 90% repolarization in a group of patients who had undergone cardioversion from either AF or AFL. More recently, Sparks et al\(^7\) demonstrated reversible decreases in lateral RA effective refractory period after brief AFL in humans and a reversal of ER after ablation of chronic AFL and resumption of SR.

In the present study, AFL produced similar changes to those observed with AF or rapid pacing, including significant decreases in AERP and CV and a significant increase in LA size. However, in contrast to atrial pacing models of AF, rapid atrial stimulation produced by an anatomic model of flutter did not usually result in spontaneous degeneration to AF but rather the stabilization of the flutter circuit. It is presumed that the presence of a long anatomic barrier and a wide EG prevented spontaneous degeneration to AF despite the ongoing rapid atrial stimulation.

When critically timed additional triggers were introduced during either SR or AFL, the ER produced by sustained AFL did provide the electrophysiological milieu for development of sustained AF that had not been present at baseline. Indeed, 28 days of AFL led to a profound increase in AF vulnerability and a gross increase in the duration of induced AF episodes from <30 seconds at baseline to >2 days. The conversion of AFL to AF was dependent on multiple (>2) tightly coupled atrial extrastimuli interacting with the AFL circuit. These extrastimuli are possibly analogous to focal triggers that arise in the pulmonary veins, associated with the initiation of focal AF or conversion of AFL to AF.\(^9\) The explanation for the different effect of LAA and RAA extrastimuli on conversion of AFL to AF or termination of AFL is unclear. The tighter CI of LAA extrastimuli, greater distance of the LAA stimulation site from the AFL circuit, and multiple potential LA-to-RA wave-front breakthrough sites\(^26\) may have resulted in a wave front that was more disorganized and fractionated when it encountered the AFL circuit.

Although an increase in AERP\(_{\text{SEP}}\) was not observed in the present study, the pattern of change (early fall then plateau) is similar to that described by Wijffels and coworkers.\(^5\) Differences between these observations and the results of other published studies may reflect either interspecies differences or variation in the distribution and density of sites chosen for AERP sampling.

**Study Limitations**

The relatively small number of chronic electrodes sutured onto the atria prevented a detailed assessment of the heterogeneity of AERP and CV changes and an exact definition of the AFL circuit. However, we did not set out to define the precise mechanisms by which AFL may cause the substrate for AF but rather to prospectively assess the stability of chronic AFL and the interaction between substrate and triggers. Furthermore, the reentrant AFL circuit generated by this Y-lesion model has been investigated extensively by the original authors.

**Conclusions**

In this anatomic model of chronic AFL, progression to sustained AF was dependent on the interaction between the development of suitable atrial substrate and the presence of focal triggers or initiators. These findings may have implications for the relationship between AFL and AF in humans.

**Acknowledgment**

Dr Morton is funded by a National Heart Foundation of Australia postgraduate medical research scholarship.

**References**


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Circulation. 2002;105:258-264
doi: 10.1161/hc0202.102012

Circulation is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
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Print ISSN: 0009-7322. Online ISSN: 1524-4539

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