Chapter from the book *Cardiac Pacemakers - Biological Aspects, Clinical Applications and Possible Complications*

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Cardiac Resynchronization Therapy: 
Lead Positioning and Technical Advances

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1. Introduction

Cardiac resynchronization therapy (CRT) is a therapeutic option for heart failure patients with a severely reduced left ventricular ejection fraction and left bundle branch block (Cleland et al., 2001). Ventricular resynchronization is achieved by biventricular pacing, usually via electrodes in the right ventricular apex and a left ventricular (LV) electrode positioned in a coronary vein. About one third of implanted patients do not respond to CRT (Derval et al., 2010). In order to reduce the percentage of non-responders, several strategies have been developed. They include optimization of patient selection, device programming as well as LV lead location. In cardiomyopathy with left bundle branch block, the lateral wall is the site of latest activation and should be the optimal location for LV pacing. Therefore, standard implantation sites for LV leads are lateral or posterolateral branches of the coronary sinus. Congruent to these pathophysiological findings, Butter et al. demonstrated a superiority of lateral wall pacing versus anterior wall pacing in CRT (Butter et al., 2001). However, a more detailed look at optimal pacing locations might be required to increase the effect of CRT and decrease non-responder rates. Different imaging modalities have been used to both identify optimal pacing sites as well as to plan LV lead implantation.

2. Imaging for cardiac resynchronization therapy

Imaging for CRT is focused on imaging of the coronary venous (CS) system for CS lead implantation and on imaging techniques to assess the left ventricular function for patient selection, choose the optimal lead position and to evaluate the effect of CRT. Contrast angiography is commonly used for imaging of the coronary venous system. To evaluate ventricular function including dyssynchrony, transthoracic echocardiography is commonly applied as it is widely available and inexpensive. A lot of efforts have been done to improve patient selection by echoardiographic screening and there are hundreds of papers published on echocardiographic evaluation of mechanical dyssynchrony, including the use of tissue Doppler imaging, speckle tracking, three-dimensional and contrast
echocardiography. However, in the PROSPECT trial with almost 500 patients no single echocardiographic parameter could predict response with convincing sensitivity and specificity (Chung et al., 2008). Despite good results in single-center studies, echocardiography for assessment of dyssynchrony is limited by high intra- and inter-observer variability, measurement errors and in some patients low image quality. Alternatives to echocardiography include magnetic resonance imaging, computed tomography and nuclear imaging. Magnetic resonance imaging has the benefit of high spatial resolution, high reproducibility and information on viability. A high scar burden and pacing over a posterolateral scar are associated with poor response to CRT (Bleeker et al., 2006, White et al., 2006). Whereas magnetic resonance imaging is the gold standard to assess myocardial viability, computed tomography also provides information on scar burden and localization as well as left ventricular function and dyssynchrony. However, data on dyssynchrony measured by computed tomography are limited and there are no published data for the prediction of CRT response. In addition, computed tomography is associated with radiation exposure. Nuclear imaging with single photon computed tomography and positron emission tomography is also associated with radiation exposure. Nuclear imaging provides information on scar burden and scar localization, ventricular function and dyssynchrony. However, a major disadvantage of nuclear imaging is the low spatial resolution.

2.1 Imaging of the coronary venous system

Left ventricular leads are usually implanted in a lateral or posterolateral branch of the coronary sinus. Contrast venography is a standard procedure performed either before or during implantation to identify suitable target veins. Table 1 displays imaging modalities for the coronary venous system.

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<th>Advantages</th>
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<td>Retrograde contrast CS angiography</td>
<td>Good vessel visibility Can be performed during CRT implantation</td>
<td>invasive</td>
<td>standard procedure</td>
</tr>
<tr>
<td>Rotational CS angiography</td>
<td>3D imaging Can be performed during CRT implantation</td>
<td>invasive</td>
<td>limited experience</td>
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<td>Computed tomography</td>
<td>Non-invasive radiation exposure</td>
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<td>MRI</td>
<td>Non-invasive lower spatial resolution</td>
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<tr>
<td>Venous phase CS angiography</td>
<td>Can be performed during standard coronary angiography</td>
<td>lower vessel visibility</td>
<td>limited experience</td>
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Table 1. Imaging modalities for the coronary venous system

Because a lot of patients with heart failure undergo a coronary angiogram, we compared retrograde occlusion venography with venous phase imaging of the coronary sinus in 24 patients (Mischke et al., 2007).
Suitable target vessels for LV lead implantation were identified in all patients by both imaging modalities. Although visibility was superior in retrograde venography than in venous phase imaging, this technique might be an alternative to retrograde venography in patients undergoing a coronary angiogram. Venous phase angiography is time-saving and easy to perform. Figures 1 and 2 display a venous phase coronary sinus angiography and a retrograde occlusion venography.

Some of the standard C-arms used for fluoroscopy allow rotational angiography with a 3D image. In an animal model we compared rotational coronary sinus angiography to ECG-gated enhanced cardiac dual source computed tomography (Knackstedt et al., 2008a). We found no significant difference between these imaging modalities with respect to vessel
diameters or vessel visibility. In contrast to computed tomography, rotational angiography images can be obtained within seconds in the catheter lab without a time lag between cardiac imaging and procedure. In addition, estimated radiation exposure and the amount of contrast medium were lower in rotational coronary sinus angiography.

We compared retrograde coronary sinus angiography with multi-slice computed tomography for visualization of the coronary venous system in 20 patients with congestive heart failure (Knackstedt et al., 2008b). Vessel visualization was better using retrograde coronary sinus angiography except for the middle cardiac vein and small veins, which were better seen with computed tomography. There was a trend that computed tomography detected more vessels. Overall, retrograde coronary sinus angiography offered a better display of target vessels commonly used for LV lead implantation.

Imaging of the coronary sinus via cardiac computed tomography correlates well with direct coronary sinus venography (Van de Verie et al., 2006). Imaging of the coronary venous anatomy by computed tomography is non-invasive and can help to plan coronary sinus lead implantation especially in patients with angulated coronary veins and in patients in whom a left marginal vein or a posterior vein is absent. It also provides information on the localization of the left phrenic nerve in relation to the target vein. However, it is associated with radiation and contrast agent exposure.

Magnetic resonance imaging of the coronary venous anatomy allows adequate assessment of localization, size and angulations of the veins (Ma et al., 2010; Nezafat et al., 2007). However, spatial resolution is inferior to computed tomography and magnetic resonance imaging cannot routinely be performed in patients with cardiac implants, e.g. patients scheduled for an upgrade from an implantable cardioverter defibrillator (ICD) to a CRT device.

2.2 Magnetic resonance imaging in CRT patients

Implanted devices like pacemakers and ICDs generally pose a contraindication to magnetic resonance imaging. However, magnetic resonance imaging might be performed with only small risks for the patient and device. In order to reduce the risks, careful patient selection, constant monitoring, specific absorption rate management and careful device programming before the scan have been used in the past. In a recent study Wilkoff et al. evaluated the safety of a pacemaker especially designed for safe magnetic resonance imaging (Wilkoff et al., 2011). Several modifications were used to improve the safety of magnetic resonance imaging, including modification of the leads to reduce lead tip heating, reduction of the amount of ferromagnetic materials, replacement of the reed switch by a Hall sensor, whose behaviour in a static magnetic field is predictable. In this prospective randomized study magnetic resonance imaging with a 1.5 T scanner could be performed without adverse events.

3. Left ventricular lead implantation

Cardiac resynchronization requires left ventricular pacing. The standard approach for left ventricular lead implantation is a transvenous implantation into a lateral or posterolateral tributary of the coronary sinus. Although dedicated instruments allow successful LV lead implantation in most patients, failure rates of 5-17% have been reported (Abraham et al., 2002; Al-Khadra et al., 2005; Purrfellner et al., 2000). In these patients, LV leads are usually implanted onto the LV epicardium through thoracotomy or thoracoscopy. In addition, there are several alternatives to standard LV lead implantation techniques, including endocardial LV lead implantation, LV lead implantation assisted by magnetic navigation and video-
assisted pericardioscopic epicardial LV lead implantation. As the access to LV regions is limited by the anatomy of the venous coronary system in standard procedures with transvenous CS lead implantation, some strategies aim at improved access to LV regions:

- Epicardial stimulation
- Endocardial LV stimulation
- Magnetic navigation for CS lead implantation
- Microcatheter LV stimulation

Table 2 lists advantages and risks of LV lead implantation techniques.

<table>
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<tr>
<th>Technique</th>
<th>Advantages</th>
<th>Disadvantages</th>
<th>Clinical relevance</th>
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<tr>
<td>Standard transvenous implantation into CS tributary</td>
<td>Low periprocedural risk</td>
<td>Limited access to LV regions, Risk of dislocation, Failure in 5-17%</td>
<td>Standard implantation procedure</td>
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<tr>
<td>Epicardial implantation via thoracotomy or thoracoscopy</td>
<td>Access to all LV regions</td>
<td>Surgical risks</td>
<td>First-line alternative to transvenous LV lead implantation</td>
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<td>Endocardial LV lead implantation</td>
<td>Access to all LV regions, Fast impulse propagation</td>
<td>Risk of thromboembolism</td>
<td>Limited experience with patients</td>
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<tr>
<td>Magnetically navigated CS lead implantation</td>
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<td>Video-assisted pericardioscopic epicardial implantation</td>
<td>Access to all LV regions</td>
<td></td>
<td>Experimental</td>
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Table 2. LV lead implantation techniques

3.1 Alternatives to right ventricular pacing

Whereas right ventricular apical pacing is the standard for patients requiring a pacemaker, this mode of stimulation is associated with electromechanical dyssynchrony and may contribute to worsening of the cardiac function (Tantengco et al., 2001). Because of the detrimental effects of right ventricular apical pacing several strategies have been suggested to avoid or reduce right ventricular apical pacing, including biventricular pacing either by de novo implantation of a CRT device or by upgrading an existing pacemaker, changes in programming to reduce the percentage of right ventricular pacing and alternative pacing sites. Careful patient selection and minimal ventricular pacing algorithms can substantially reduce the amount of right ventricular pacing and have been implemented into clinical practice (Tops et al., 2009). Several studies have demonstrated a hemodynamic and symptomatic benefit of upgrading right ventricular apical pacing to CRT as well as CRT in patient with indications for permanent pacing (Tops et al, 2009). However, so far it remains uncertain whether this will translate into a prognostic benefit. Alternative pacing sites have been suggested to avoid right ventricular apical pacing, including pacing the right
ventricular outflow tract, septal pacing and direct His bundle pacing. A meta-analysis by de Cock (de Cock et al., 2003) showed a favorable hemodynamic effect, and a study by Venerio (Venerio et al., 2008) demonstrated an improved survival in patients with right ventricular outflow tract pacing as compared to right ventricular apical pacing. However, most studies include rather small numbers of patients and are of short follow up, so more data are needed to evaluate the relevance of right ventricular outflow tract pacing for clinical routine. Septal pacing has been shown to decrease ventricular dyssynchrony compared to right ventricular apical pacing (Yu et al., 2007) but there was no difference in left ventricular ejection fraction in a prospective study by Kypta (Kypta et al., 2008). Direct His-bundle pacing or para-Hisian pacing allows a more physiological impulse propagation than right ventricular apical pacing but is associated with difficulties in lead positioning and concerns about pacing thresholds (Tops et al., 2009).

Henz (Henz et al., 2009) demonstrated in a small animal study the feasibility of atrioventricular septal synchronous pacing with intramyocardial leads implanted deep within the atrioventricular septum; further animal studies are needed to evaluate this approach.

### 3.2 Optimized CS lead implantation

Already a decade ago Butter et al. demonstrated a hemodynamic superiority of pacing from a lateral vein compared to an anterior vein for CRT (Butter et al., 2001). The distance between stimulation site and the region of latest contraction may be crucial for hemodynamic benefit of CRT (Ypenburg et al., 2008). This is in line with findings from animal studies (Helm et al., 2007) and studies using echocardiographic parameters in patients (Becker et al., 2007a and 2007b).

We used computed tomography and MRI imaging prior to LV lead implantation in 20 patients with congestive heart failure (Knackstedt et al., 2010a). Computed tomography was used for imaging of the coronary venous system and MRI to detect the region of latest contraction. Computed tomography and MRI images were then over-imposed to determine a coronary side branch suitable for lead implantation that is closest to the region of latest contraction. There was a trend towards a shorter distance between the LV lead and the region of latest contraction in patients classified as responders.

Another approach is the use of myocardial deformation analysis assessed by circumferential strain analysis during echocardiography to determine the optimal site for CS lead implantation. In a study with 56 patients optimal LV lead position was defined as a lead position close to the segment with latest systolic strain prior to CRT (Becker et al., 2010). During follow up, patients with leads implanted in an “optimal position” experienced a significantly higher increase in left ventricular ejection fraction than patients with leads implanted at other sites.

In a smaller study Ducket et al. performed computed tomography and MRI to acquire 3D whole heart images. After segmentation, 3D anatomical models were overlaid over live fluoroscopy to guide LV lead implantation (Ducket et al., 2010).

### 3.3 Endocardial LV lead implantation

Endocardial lead implantation is associated with a high risk of systemic thromboembolism (van Gelder et al., 2000). However, endocardial LV lead implantation has several (potential) advantages to CS and epicardial LV leads: it allows access to all LV regions, endocardial ventricular layers offer faster impulse propagation than epicardial layers and endocardial stimulation might result in improved hemodynamics. Van Deursen demonstrated in an
acute canine model a superior electrical resynchronization as well as $+dP/dT_{\text{max}}$ when endocardial biventricular stimulation was used instead of epicardial stimulation. In addition, whereas epicardial stimulation resulted in a transmural dispersion of repolarization, this was not observed in endocardial stimulation (van Deursen et al., 2009). However, Spragg et al. compared the hemodynamic effects of endocardial pacing at sites directly transmural to the CS lead tip in a small study of patients and found no difference in hemodynamics (Spragg et al., 2010). In this study a superior hemodynamic result was seen in 8 of 11 patients when endocardial pacing was performed from extreme basal sites at positions adjacent to the mitral ring. In a study by Derval et al. (Derval et al., 2010) pacing at the best LV site in 35 patients with non-ischemic dilated cardiomyopathy was associated with twice the improvement in $+dP/dT_{\text{max}}$ compared to CS pacing.

In summary, the major benefit of endocardial left ventricular pacing seems to be the access to all LV regions, whereas endocardial stimulation per se seems to be only of minor relevance.

In a few patients with major surgical contraindications to epicardial LV leads have been implanted through a transseptal approach (Jais et al., 2000; Leclercq et al., 1999; van Gelder et al., 2007). However, this approach is technically challenging. In addition to the risk of thromboembolism due to leads in the LV cavity, the adjacency to the mitral valve carries the risk of mitral insufficiency as well as endocarditis in case of infectious complications. A transapical approach which has been described by Kassai et al. in a limited number of patients would avoid passage of the mitral valve (Kassai et al., 2008).

### 3.4 Magnetically navigated LV lead implantation

A tortuous course of the coronary venous tree and target veins with small diameters can sometimes be challenging for CS lead implantation. New wire and lead navigation systems might facilitate lead implantation. The Niobe System (Stereotaxis Inc., St. Louis, USA) allows remote magnet controlled navigation of catheters and guidewires. The magnetically navigation system consists of two permanent magnets creating a steerable magnetic field (figure 3). The magnetic guidewires include a small magnet at their tip and can be steered by changing the orientation of the outer magnets. The magnetic field vector is displayed on a monitor and can be changed from the control room or from a bedside touch-screen monitor with sterile covers (figure 4).

We studied 123 patients who were assigned to either conventional CS lead implantation or LV lead implantation using magnetic navigation (Mischke et al., 2009). Venography of the coronary venous system was performed to select a target vessel for lead implantation. Left ventricular lead placement was analyzed with regard to three endpoints: 1) engagement of the target vessel with the guidewire, 2) over-the-wire lead placement in the target vessel, and 3) final LV lead position. Guidewire access to the target vessel was achieved in all patients using magnetic navigation compared to 87% with the conventional approach ($p<0.05$). Implantation success rates, total procedure and fluoroscopy times did not differ significantly between groups. Gallagher et al. used the Niobe system for CRT implantation in 50 patients (Gallagher et al., 2007). In this study, vessels were engaged either by CS venography and the use of a magnetic guidewire or via a “bare wire” approach without venography or special CS delivery sheaths. For the “bare wire” approach, the guidewire was used to probe for a target vessel as a substitute for CS venography. This was associated with a reduction in procedure and fluoroscopy time compared to the use of CS sheaths and venography.
Fig. 3. Catheter suite with a magnetic navigation system (Niobe). Permanent magnets to both sides of the fluoroscopy table can be moved inside their casings to alter the magnetic field.

Magnetic navigation might be used as an additional tool for precise wire navigation and enable the operator to engage target vessels that are tortuous. In addition, technical advances in lead design might allow engagement of vessels which are now being considered inadequate due to morphology or size. We have recently demonstrated the feasibility of left ventricular stimulation via a miniaturized magnetized stimulation wire in an acute animal model (Knackstedt et al., 2010b). A conventional guide wire with a permanent magnet and a single stimulation electrode at its tip was coated with iridium oxide at the distal end and insulated except for the very tip. The stimulation wire was steered into side branches of the coronary sinus via magnetic navigation and successful left ventricular stimulation was performed via the wire.

Fig. 4. Navigant screenshot. Two x-ray images have been transferred to the navigation software and tributaries of the CS have been marked with colors. The red arrow indicates the direction of the magnetic field vector.
3.5 Video-assisted pericardioscopic epicardial lead implantation
In an animal study we used flexible and rigid endoscopy for implantation of epicardial pacing leads via a subxiphoidal access (Hatam et al., 2010). Rigid endoscopy showed to be superior to flexible endoscopy with regard to stability and orientation within the pericardial space, and leads were successfully implanted onto all four cardiac chambers. This minimally invasive procedure allows access to all left ventricular regions. However, this technique requires a subxiphoidal access to the epicardial space and the endoscopy is associated with ventricular arrhythmias.

3.6 Leadless pacing
Pacing without pacemaker leads would decrease the risk of infection and might allow multisite pacing and thus decrease dyssynchrony. Ultrasound and magnetic field waves have been used to induce electrical stimulation via an intracardiac receiver electrode (Kapa et al., 2010; Lee et al., 2007). In an animal study Echt et al. (Echt et al., 2006) used burst ultrasound energy transmission through the chest to a receiver electrode mounted on a catheter that converted the ultrasound energy to electrical energy sufficient to pace the myocardium. Biventricular pacing was also possible in this acute animal study. Microscopic evaluation revealed no evidence of mechanical or thermal bioeffects. Lee et al. successfully tested this system in patients undergoing electrophysiological studies (Lee et al., 2006). The technology in this study is under development as a leadless implantable system for chronic use. Technical challenges include a high beat-to-beat variation in the receiver electrode output as well as inefficient energy conversion: less than 1% of the transmitted energy was used for cardiac pacing.

In an acute animal study Wienke (Wieneke et al., 2009) demonstrated the feasibility of cardiac pacing via induction technology. The systems consisted of a transcutaneously implanted transmitter unit made of a ring-shaped copper coil and a receiver unit implanted in the right ventricular apex. The transmitter generated an alternating magnetic field of around 0.5 mT that was converted into a voltage pulse by the receiver in order to pace the ventricle. So far results have been published from one pig only, and no data on chronic pacing are available.

A promising miniaturized leadless pacemaker is being developed by Medtronic (Minneapolis, USA): the small device can be deployed with a catheter from a venous access and implanted into the ventricular cavity. Up to now no animal or human data have been published about the device.

4. Electrical remodeling in CRT
In congestive heart failure (CHF), a complete left bundle branch block causes asynchronous ventricular contraction due to regional dispersion of ventricular depolarization, resulting in intra- and interventricular mechanical asynchrony. CRT reduces the heterogeneity of ventricular contraction by biventricular stimulation.
Especially patients with a very broad QRS-complex (> 150 msec) seem to profit most from CRT (Chung et al., 2008; Moss et al., 2009). Although QRS duration is not an optimal criterion for selecting patients amenable for CRT and some studies have failed to predict clinical and echocardiographic response to CRT, it remains an important criterion for dyssynchrony for the indication of CRT (Boriani et al., 2006; Gervais et al., 2009; Hawkins et al., 2006; Mollema et al., 2007).
In CRT, biventricular stimulation usually results in a narrowing of the stimulated QRS-complex and a reduction in left ventricular chamber size as well as improvement in ejection fraction. The extent of the QRS shortening induced by biventricular pacing seems to correlate with the structural remodeling (Boriani et al., 2006; Kronborg et al., 2010). However, about one third of patients fail to respond to CRT (Cleland et al., 2001; Chung et al., 2008; Lafitte et al., 2009). A lot of effort has been spent to both identify patients who are likely to benefit from CRT and to increase the benefit from CRT, e.g. by optimizing AV and VV delays (Strauss et al., 2010).

Although CRT has been shown to induce a structural remodeling resulting in reduction in left ventricular dimensions and improvement in ejection fraction, there is scarce and controversial data on a possible remodeling of the native conduction system (Dizon et al., 2004; Henrikson et al., 2007; Stockburger et al., 2008).

We studied the effect of CRT on the native conduction system in a small prospective study (Mischke et al., 2011). A CRT device was implanted in 38 patients with congestive heart failure (ejection fraction (EF): 26 ± 7%). 20 patients suffered from dilated cardiomyopathy and 18 from ischemic cardiomyopathy. Standard 12-lead ECGs with and without pacing as well as echocardiographies were obtained prior to implantation and after 6 and 12 months. Patients were classified as responders in case of an increase in EF ≥ 25% in combination with an increase in NYHA class ≥ 1. The EF increased to 36 ± 10% (p<0.0001) after 6 months and 40 ± 12% (p<0.0001) after 12 months of CRT. Intrinsic QRS duration decreased from 171 ± 18 ms before CRT to 164 ± 23 ms (p=0.027) after 6 months and 161 ± 25 ms (p=0.002) after 12 months of CRT (figure 5). 22 patients (58%) were classified as responders. Whereas a significant decrease in intrinsic QRS duration was observed in responders, only a slight decrease was seen in non-responders. However, two-factorial variance analyses did not show a significant influence of response or underlying heart disease (dilated or ischemic cardiomyopathy) on the change in QRS duration (p=0.7).

![Fig. 5. Baseline and intrinsic QRS duration (from Mischke et al., 2011)](image)

This was the first prospective study to demonstrate a decrease in intrinsic QRS duration in patients treated with CRT. Dizon et al. reported the first case of loss of bundle branch block in a patient 6 months after implantation of a CRT device (Dizon et al., 2004). However, data on intrinsic QRS duration is controversial. No change in intrinsic QRS duration was seen in
the MUSTIC trial as well as in a study by Stockburger (Stockburger et al., 2008). Two studies displayed a trend towards a reduction in intrinsic QRS duration (Boriani et al., 2006; Vogt et al., 2000). Similar to our results, a retrospective study by Henrikson et al. showed a significant reduction in intrinsic QRS duration after 14 months of CRT in 25 patients (Henrikson et al., 2007). Experimental data suggest a subcellular redistribution of connexin43 and ion channel remodeling with a reduction in inward rectifier K⁺ current, delayed rectifier K⁺ current and transient outward K⁺ current) and abnormal Ca²⁺ homeostasis in left bundle branch block (Aiba et al., 2009; Spragg et al., 2005). CRT partially restored this ion channel remodeling and attenuated the regional heterogeneity of action potential duration. Although human data on intrinsic QRS duration in CRT is controversial, an impact on the conduction system by several factors including connexin redistribution and reduction in left ventricular dimensions is quite conceivable.

5. Conclusion

Cardiac resynchronization therapy is an effective treatment for patients with congestive heart failure and complete left bundle branch block. However, about one third of all patients who undergo CRT do not profit from it. Several strategies have been tried to reduce the percentage of non-responders, including optimized patient selection, device programming and optimized positioning of the left ventricular lead. However, due to high interpatient variability there seems to be no single best pacing site for all patients. Acute hemodynamic testing during implantation is time-consuming and good acute effects might not translate into a long-term clinical benefit.

None of these approaches has had a relevant impact on daily practice yet. In order to have the maximum benefit for our patients, we need to individualize the approach to CRT. Technical advances, like new lead designs and guiding catheters, are crucial for the further progress in CRT.

6. References


Clinical usage of artificial pacing dates back to 1958, when the battery powered cardiac pacemakers became available. Modern implantable pacemakers are the complicated electronic devices operating 10 years continuously without battery exchange. Though the development of devices is not a primary topic of the book, certain efforts towards developing of biologic pacemakers through tissue engineering and studying of cell synchronization are discussed. The main attention is paid to implementations of pacemakers in different medical situations oriented towards widening the clinical indications for implanting the cardiac pacemakers. New methods and devices in cardiac resynchronization therapy (CRT) have received particular attention. Placing of pacing electrodes has been treated soundly. Furthermore, emerging of complexities and complications in new clinical situations and other safety problems have been discussed thoroughly. The authors have derived the used information from their own clinical practice and experiences of their medical colleagues. These and other pragmatic features can be acknowledged as the most valuable asset of the book.