A study of ECG pattern, cardiac structural abnormalities and familial tendency in patients with early repolarisation syndrome in South India

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ABSTRACT

Background Early repolarisation (ER) on ECG, which was initially believed to be benign, has of late been considered otherwise. Brugada syndrome has recently been thought to be an extension of the ER spectrum, and the familial tendency of the ER pattern is being highlighted. With attention being drawn to ER’s association with idiopathic ventricular fibrillation (VF), the prognosis and lineage of patients with an ER pattern are under scrutiny.

Aims To analyse ER patterns on ECG, their presence in first-degree relatives and their association with structural heart disease. To classify different types of ER and estimate the prevalence of the high-risk notch/slur pattern in the population studied.

Methods We screened all patients presenting to our department from December 2011 to July 2014 for ER patterns. We excluded patients with other causes of ST elevation that mimicked the ER pattern, those aged <18 years, and those not willing to participate in the study. A complete physical examination, 12-lead ECG and echocardiography were performed on all study patients. Willing first-degree relatives were screened with a 12-lead ECG. Of the 963 patients with ER that we initially screened, 843 completed the study. A total of 4116 relatives were screened.

Results Of the 843 patients who completed the study, 687 (81.5%) were male and 156 (18.5%) were female. The majority were asymptomatic (70.11%), but had been referred for ECG abnormalities. Fifteen patients with chest pain were inadvertently thrombolysed and were later diagnosed to have ER. Their ER pattern was exaggerated during chest pain, which made this error highly likely. Among the 48 patients who had acute coronary syndrome (ACS), ER pattern was noticed in a different lead than those affected by ACS. Of these, 27 (56.25%) had ventricular tachycardia/VF during the acute phase. Six patients had electrical storm without evidence of ACS, and all had a global ER pattern with prominent notching/slurring on baseline ECG. The most common type of ER pattern was type I (lateral leads; 55.87%). Twenty-one patients had a Brugada pattern. Of all the patients with ER, only a third (34.16%) had the possibly high-risk notched/slurred ECG pattern. The majority (82.92%) had a structurally normal heart. We found that mitral valve prolapse (MVP), as assessed by >2 mm leaflet prolapse from the annulus, was more common in patients with ER (11.39%). Of the 4116 relatives screened, 2625 (63.78%) had an ER pattern; a quarter of family members had the same ER pattern. We also noticed different ER patterns in the same family.

Conclusions We found that exaggeration of the ER pattern during chest pain may lead to inadvertent thrombolysis. A notched/slurred ER pattern is found in only a third of patients, who need to be grouped separately, as they may constitute a high-risk category. Patients with ER had MVP at a higher prevalence (almost double) than the general population, probably explaining the high incidence of sudden cardiac death associated with MVP. A familial tendency to an ER pattern was found in more than half of first-degree relatives, with different ER patterns, even the Brugada pattern, found in the same family. This may be because Brugada and other ER patterns belong to the same spectrum and may share the same prognosis. Thus we conclude that further studies regarding ER, its association with MVP, risk stratification by notched ECG pattern, and familial distribution along with gene analysis are warranted.

BACKGROUND

Early repolarisation (ER) on ECG was detected as early as 1936 and was considered benign for a long time. In 1984, the malignant nature of ER was first noted and was confirmed by several studies, including the seminal paper of Haïssaguerre et al on ER’s association with idiopathic ventricular fibrillation (VF). Brugada syndrome has been proposed to be an extension of the ER pattern and has become an indication for an implantable cardioverter defibrillator. Bearing this fact in mind and considering that ER has a familial tendency, we analysed ECG patterns of patients with ER presenting to our hospital, along with familial tendency to ER and its association with any structural heart disease. We were also inspired by the fact that ER is a current ‘hot’ topic and not many studies on ER have been reported from this part of the country.

METHODS

We analysed the ECGs of all patients presenting to the Department of Cardiology, Government T D Medical College Hospital, Alappuzha, Kerala, South India from December 2011 to July 2014 for ER change. For our study, ER was defined as ‘an ECG change characterised by J-point elevation and rapidly upsloping or normal ST segment of ≥1 mm (0.1 mV) above the isoelectric line and present in at least two contiguous leads’ and was derived from American Heart Association/American College of Cardiology Foundation/Heart Rhythm Society recommendations for the standardisation and interpretation of the ECG published in Circulation (2009). Exclusion criteria were <18 years of age, not willing to sign the informed consent form, and...
other proven causes for ST elevation. Patients whose investigations could not be completed were also excluded. The majority of patients were referrals from other departments, especially the Department of Anaesthesia, for ECG abnormalities. Of the 963 patients initially screened, 843 completed all the necessary investigations and gave informed consent. All these patients had a complete physical examination, 12-lead ECG, and echocardiography. Other investigations were performed according to relevance on a case-by-case basis. First-degree relatives also signed an informed consent form and were screened with 12-lead ECG alone. The total number of relatives screened was 4116. The results were tabulated, and relevant statistical analysis was performed.

**RESULTS**

Of the 963 patients recruited, 843 were included in the study; 687 (81.5%) were male and 156 (18.5%) were female. The mean age of the study population was 47.3 years for men and 44.7 years for women. The average heart rate was 57/min for women and 62/min for men. The vast majority were asymptomatic (70.1%) and were referred from other departments for ECG abnormalities, mainly detected during preoperative evaluation (figure 1). The next most common presentation was chest pain, the majority of which was atypical pain. Fifteen of these patients were inadvertently thrombolysed and later diagnosed with ER syndrome, as echocardiography, troponin and coronary angiography (CAG) results were negative. We also noted that the ER pattern in these patients was exaggerated during chest pain, which made this error highly likely. Forty-two patients presented with syncope, but neither their history nor ECG or Holter monitor results suggested significant arrhythmias. Forty-eight patients had acute coronary syndrome (ACS) and an ER pattern in a different lead from those affected by ACS. Of these, 27 (56.25%) had ventricular tachycardia (VT)/VF during the acute phase. However, we did not notice any association between any particular type of ER and VT/VF. Six patients had electrical storm without evidence of ACS as documented by enzyme assay or echocardiography. These patients ultimately died. We convinced the relatives of three such patients to allow an autopsy, which showed a flabby heart without evidence of ACS. All had a global ER pattern with prominent notching/slurring on the baseline ECG.

Among the ECG patterns of ER (table 1), the most common was type I—that is, involving lateral leads (55.87%). We found 21 patients with a Brugada pattern (all three types in total), which we grouped with other ER patterns, as the sample size was small. We also grouped patients with a notched/slurred ECG pattern (figure 2) separately to determine its relative prevalence in each type of ER (figure 3).

Echocardiography was performed in all patients to detect structural heart disease (table 2). The majority (82.92%) had a structurally normal heart. We found mitral valve prolapse

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**Table 1** ECG patterns in patients with early repolarisation

<table>
<thead>
<tr>
<th>Lead involved</th>
<th>No of patients (n=843)</th>
<th>Percentage</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lateral</td>
<td>471</td>
<td>55.87</td>
<td>50.06 to 61.68</td>
</tr>
<tr>
<td>Inferior</td>
<td>162</td>
<td>19.22</td>
<td>14.61 to 23.83</td>
</tr>
<tr>
<td>Inferolateral</td>
<td>141</td>
<td>16.73</td>
<td>12.37 to 21.09</td>
</tr>
<tr>
<td>Global</td>
<td>6</td>
<td>0.71</td>
<td>−0.27 to 1.69</td>
</tr>
<tr>
<td>Other patterns</td>
<td>63</td>
<td>7.47</td>
<td>4.4 to 10.54</td>
</tr>
</tbody>
</table>

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**Figure 1** Presentation of patients with early repolarisation. ACS, acute coronary syndrome; VF, ventricular fibrillation; VT, ventricular tachycardia.

**Figure 2** Examples of notched/slurred ECG patterns.
(MVP), as assessed by >2 mm leaflet prolapse from the annulus, to be more common in patients with ER (11.39%). A total of 117 patients underwent CAG, of which only 48 had significant coronary artery disease (CAD).

To estimate the familial tendency of ER patterns, we screened willing first-degree relatives of the patients with a 12-lead ECG. If at least one of the screened relatives had an ER pattern in any of the leads, it was considered a familial tendency. Of the 4116 relatives screened in total, 2625 (63.78%) had an ER pattern. A quarter of family members with the inferolateral variety and over 60% of relatives with the lateral variety had familial ER patterns (table 3 and figure 4). We also noted that the ER pattern was not necessarily the same in all family members (table 4). Of the 4116 first-degree relatives screened, 2625 had an ER pattern in one of the leads. The first column in table 4 shows the ER pattern in index cases, and the following columns depict the ER pattern found in relatives of each of the ER patterns in the first column.

**DISCUSSION**

ER syndrome is an as yet unexplored subject, and the potential danger associated with ER was only discovered recently. ER has been classified into three types. Type I has the best prognosis, with ECG changes confined to lateral leads. Type II has moderate risk and has changes confined to inferior or inferolateral leads. Type III has global ER changes and has the worst prognosis. The Brugada pattern is at present considered to be an extension of the ER pattern. Brugada syndrome has been proven to be arrhythmogenic and a definite familial relation has been established, while the three subgroups of ER have no defined management protocols.

Of the ER patients presenting to our department, the majority were asymptomatic, as all surgical patients were screened with an ECG before surgery. Most patients with chest pain had atypical pain, and screening for CAD proved negative in the majority. Fifteen patients were inadvertently thrombolysed, as there was an augmentation of ER changes mimicking ST elevation myocardial infarction during their chest pain. This augmentation just before the onset of arrhythmia is well documented. A total of 117 patients underwent CAG. Only 48 of these patients had significant CAD and were managed accordingly. We did not perform a vasoactive test to rule out vasospastic angina in any of these patients, as it was not considered essential and hence not cleared by the institutional ethics committee.

We also noticed that it is possible to have both ACS and ER changes on the same ECG but in different leads. Of the 48 patients with ACS, 27 developed VT/VF during ACS. Interestingly, all 27 had a notched/slurred QRS pattern, and 15 of them had ER changes in inferior leads. Many studies that suggested an association of idiopathic VF with ER pattern also found notching on ECGs of affected patients. This is also true with Brugada syndrome, where right bundle branch block is associated with ST elevation. The six patients in our study with idiopathic VT/VF also had a global ER pattern with notching of QRS. Thus we categorised patients with a notched ECG pattern separately to determine its relative prevalence and found that fewer than 35% of the total patients with ER had a notched ECG.

The most common type of ER pattern was type I (confined to lateral leads), and the most common population was young men. This has also been documented in other larger studies. From echocardiography results, we found the majority of

**Table 2** Echocardiographic findings in the 843 study patients

<table>
<thead>
<tr>
<th>Echo</th>
<th>Male</th>
<th>Female</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
<td>95% CI</td>
</tr>
<tr>
<td>Normal</td>
<td>585</td>
<td>85.15</td>
<td>80.54 to 89.76</td>
</tr>
<tr>
<td>MVP</td>
<td>69</td>
<td>10.04</td>
<td>6.15 to 13.93</td>
</tr>
<tr>
<td>RWMA</td>
<td>21</td>
<td>3.06</td>
<td>0.83 to 5.29</td>
</tr>
<tr>
<td>Other valves</td>
<td>12</td>
<td>1.75</td>
<td>0.05 to 3.45</td>
</tr>
<tr>
<td>Total</td>
<td>687</td>
<td>156</td>
<td>843</td>
</tr>
</tbody>
</table>

MVP, mitral valve prolapse; RWMA, regional wall motion abnormality.

**Table 3** Familial pattern of ER in relatives of index ER patients (cumulative of all patterns)

<table>
<thead>
<tr>
<th>Pattern</th>
<th>Patients with familial pattern</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lateral</td>
<td>285/471</td>
<td>60.51</td>
</tr>
<tr>
<td>Inferior</td>
<td>84/162</td>
<td>51.85</td>
</tr>
<tr>
<td>Inferolateral</td>
<td>36/141</td>
<td>25.53</td>
</tr>
<tr>
<td>Other patterns</td>
<td>33/69</td>
<td>47.83</td>
</tr>
<tr>
<td>Total</td>
<td>438/843</td>
<td>51.96</td>
</tr>
</tbody>
</table>

The number of patients whose relatives were screened for ER and at least one of them had an ER pattern in any of the leads is shown.
patients to have a structurally normal heart, as expected. However, the most interesting finding was that MVP was associated with ER (11.39%) at a higher prevalence than in the general population\textsuperscript{16–17} (maximum of 5%). This probably explains the higher incidence of palpitations and SCD associated with MVP\textsuperscript{15–17}. Further studies in this regard may help to find a definite and clear association.

However, the most interesting part of our study was the magnitude of familial tendency in these patients. Around 51.96% had an ER pattern running in the family. The most common was a lateral pattern and the least common an inferolateral pattern. Another finding was that the same family can harbour different ER patterns. Patients with a lateral ER pattern on ECG had relatives with an inferior, inferolateral or even a Brugada pattern. This may point to the fact that at least some of the ER patterns may in fact be a continuum of Brugada syndrome and may share the same prognosis. This needs further study.

**CONCLUSION**

The shift in concept of ER from being benign to being malignant has revived interest in ER. We noticed that exaggeration of the ER pattern during chest pain in certain patients may mimic ACS, which may lead to inadvertent thrombolyis. These patients usually do not have reciprocal changes in other leads, and echocardiography and troponin investigations will be negative. Patients with a notch/slur ER pattern need to be grouped separately, as they may constitute a high-risk category. Further studies may clarify this, which may help in risk stratification of patients with ER. Another interesting finding was the higher prevalence of MVP (almost double) in patients with ER than in the general population. This probably explains the higher incidence of SCD in MVP. Familial tendency of an ER pattern was found in more than half of the first-degree relatives. The most interesting point was that the ER pattern could vary in the same family, and a Brugada pattern could even be found alongside other patterns. Consequently, the possibility of Brugada and other ER patterns belonging to the same spectrum has to be considered and may even share a similar prognosis. Hence this study lays the foundation for further population-based studies on ER, its association with MVP, risk stratification by notch/slur ECG pattern, genetic screening and familial distribution.

**Collaborators** Dr Issac V Mamman, Dr Saritha S Nair, Dr MG Divakaran, Dr Gagan Velayudhan, Dr Teffy Jose, Dr Sunitha Viswanathan.

**Contributors** KGM was the primary investigator and coordinated the whole study, collected the majority of the data, and conducted the analysis. VG, TGB, RR and SK helped with collection of data, while RB, KSM, RJ and VVR helped with analysis. Other collaborators helped KGM with the whole study, as they are part of the department but to a lesser extent than the contributing authors.

**Competing interests** None.

**Patient consent** Obtained.

**Ethics approval** Institutional Review Board, Government TD Medical College, Alappuzha.

**Provenance and peer review** Not commissioned; externally peer reviewed.

**REFERENCES**


**Table 4** Familial tendency in patients with early repolarisation with respect to lead

<table>
<thead>
<tr>
<th>Lead</th>
<th>Lateral</th>
<th>Inferior</th>
<th>Inferolateral</th>
<th>Other</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
<td>n</td>
<td>n</td>
<td></td>
</tr>
<tr>
<td>Lateral</td>
<td>909</td>
<td>61.84</td>
<td>198</td>
<td>13.47</td>
<td></td>
</tr>
<tr>
<td>Inferior</td>
<td>69</td>
<td>13.07</td>
<td>285</td>
<td>53.98</td>
<td></td>
</tr>
<tr>
<td>Inferolateral</td>
<td>81</td>
<td>19.29</td>
<td>54</td>
<td>12.86</td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>36</td>
<td>17.39</td>
<td>27</td>
<td>13.04</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>1095</td>
<td>564</td>
<td>813</td>
<td>153</td>
<td>2625</td>
</tr>
</tbody>
</table>

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