Late gadolinium enhancement (LGE) progresses with right ventricle volume in children after repair of tetralogy of fallot☆☆

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1. Introduction

Congenital heart defect is present in 0.5–1% of newborn infants and 3–5% of the patients are born with tetralogy of Fallot (TOF) [1–4]. The outcome of these patients has improved dramatically during the last decades but the long-term prognosis is still compromised by important chronic sequelae like right ventricular (RV) scarring, dysfunction, dilation and arrhythmias leading to higher morbidity and mortality than in their peers [4,5]. Surgical correction of right ventricular outflow tract (RVOT) obstruction often results in pulmonary regurgitation, N-terminal pro-brain natriuretic peptide (NT-proBNP) or the aminoterminal propeptide of type III procollagen (PIIINP). We also studied if the patient’s age, post-operative follow-up time or surgical history would affect LGE.

Methods: A total of 40 pediatric patients who had undergone TOF repair and 43 healthy age and gender matched controls underwent a CMR study, whereby LGE was scored in the right (RV) and the left ventricle. To exclude the possible iatrogenic scarring we calculated the LGE score by excluding the right ventricular outflow tract and VSD patch region.

Results: All patients had RV LGE and in 39 of 40 it was seen also outside the surgically affected areas. The amount of LGE correlated positively with the RV end-diastolic volume (r = 0.44, P = 0.0045), pulmonary regurgitation (r = 0.40, P = 0.013), and with NT-proBNP. The presence of LGE also depended on post-operative follow-up time (r = 0.53, P = 0.006). PIIINP levels of TOF patients were significantly higher than in the control subjects but it did not correlate with LGE or with any of the studied clinical markers.

Conclusions: LGE is present globally in the right ventricular muscle in children and adolescents with TOF. The longer the follow-up time the more common was the LGE in the right ventricle.

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time or patient age. In addition, we studied whether LGE correlated with plasma N-terminal pro-brain natriuretic peptide (NT-proBNP), a well-established marker for heart failure [17–19], found elevated with increased RV volume and pulmonary regurgitation in adult and adolescents series of patients [20–22]. Finally, since LGE has previously been demonstrated in processes with increasing fibrosis [13–15,24] we studied whether aminoterminal propeptide of the type III procollagen (PIIINP), a biochemical marker of this fibrillar collagen turnover, would correlate with right ventricular LGE, its end diastolic volume, or the extent of the pulmonary regurgitation.

2. Methods

Forty-five patients who had undergone surgical repair of TOF between 1990 and 2003 were recruited to the study. 39 patients arrived from the Helsinki University Hospital district, serving a population base of 1.7 million. The inclusion criteria were suspicion of a severe pulmonary regurgitation in echocardiography. Six additional patients arrived from other hospitals referred by their own pediatric cardiologist according to the same criteria. Three patients declined from the study and two patients were rejected because of insufficient MRI image quality. Accordingly, forty patients were admitted to the study and were examined during an annual ambulatory visit to the clinic. For control subjects, 43 healthy age and gender matched pediatric and adolescent volunteers were recruited around the Helsinki area. They were accepted to the study if they had no medical history of any cardiovascular disease and no other pre-existing condition that would affect the cardiovascular system. No subjects had any other known conditions associated with elevated levels of plasma concentrations of PIIINP such as pulmonary fibrosis, chronic liver disease or renal dysfunction. All of the patients and control subjects underwent a clinical examination by the pediatric cardiologist. The study protocol included an echocardiogram, a CMR study, and a blood sample, from each of the subjects.

The Ethics Committee of the Children’s Hospital of the Helsinki University Central Hospital approved the study. A written informed consent was obtained from all the participants and/or their parents.

CMR images were obtained with a 1.5 T Philips Achieva imager (Philips Healthcare, The Netherlands) and a 5 channel cardiac coil. For right-ventricular-volume analysis, transaxial and left ventricle (LV) short axis balanced turbo field echo (B TFE) breath-hold cine images were obtained throughout the whole heart. Slice thickness was 5–8 mm depending on the patient size, gap was 20% and temporal resolution was 26–43 ms. For RV end diastolic volume measurements, both the transaxial and the LV short axis cine images were manually planimetered with the cardiac analysis tool of a Philips ViewForum workstation (Philips Healthcare, The Netherlands). Pulmonary arterial flow and regurgitation were assessed using a through plane phase contrast measurement. The phase contrast plane was positioned perpendicular to the pulmonary artery flow one cm above the valve. A retrospectively gated free breathing acquisition with three averages lasting typically for one minute was obtained with a temporal resolution of 24 ms and VENC 200. The field of view was kept as small as possible for optimal spatial resolution and slice thickness = 10 mm, TR = 4,019, TE = 2,427, and ET = 4. The flow images were analyzed with the workstation provided by the MR scanner vendor. A radiologist experienced in CMR performed all CMR analysis.

For LGE analysis, transaxial and sagittal late enhancement images were obtained from both ventricles after 10 min of Gd-DOTA, Gadoterate meglumine 279.3 mg/ml (Dotarem®) 0.2 ml/kg injection. A gradient echo sequence with TR = 3.239 ms, TE = 1.628 ms and ET = 49 was used. Slice thickness was 7 mm with no gap. The inversion time (TI) was determined with the use of TI-scout sequence and varied between 260 and 360 ms.

RV LGE grading protocol with seven RV segments including the target cites of repair served as a basis for our grading method [13]. For the present work we performed scoring of LGE similarly to this earlier protocol, but adapted the method by omitting 2 surgically manipulated segments (VSD patch region and the anterior wall of RVOT) to quantify remote LGE within the RVs. Accordingly, 5 remote segments were used in the analysis: anterior wall of RV, inferior wall of RV, RV surface of septum, trabecular bands and RV insertion points. The late enhancement images were visually assessed and each segment was graded according to the linear extent of enhanced myocardium. The protocol is described in detail in Fig. 1. Meticulous care was taken to confirm RV LGE from different views. The radiologist was blinded to the clinical status and other findings of the patients. Typical late enhancement findings are shown in Figs. 2 and 3. For the LV LGE scoring, a standard 17-segment model was used [23].

The radiologist blinded to the primary results later repeated scoring of 23 patients’ LGE images. A good correlation (R2 = 0.77) was found between the two assessments. Bland–Altman method was used to study intra-observer variability and found a difference in average score of 0.47. Difference of the individual scores was between −1 and +1 in 95% of the measurements.

Blood samples from a peripheral vein were drawn from all the subjects into vials with an appropriate anticoagulant agent followed by plasma separation by centrifugation. NT-proBNP was measured from the plasma by immunochemiluminometric assay at the biochemical laboratory immediately after the sample was taken. Another part of the separated plasma was stored at −70 °C for later analysis, from which plasma levels of PIIINP were measured using a commercially available radioimmunological method (Orion Diagnostica UniQ PIIINP RIA, Orion Corporation Orion Diagnostica, Espoo, Finland). All analyses were performed in duplicate, and the average was used as a single datum. The analytic data of the control patients demonstrated that circulating PIIINP did not change between 7 and 19 years of age. Accordingly, no age-dependent correction was needed to the analytic data on PIIINP.

Statistical analysis was performed using GraphPad Prism 5.0a (GraphPad Software, Inc. San Diego, California, USA). All results are reported as a mean ± SD. Comparisons of continuous data between the groups were performed using an unpaired t test. Correlations were calculated by using linear regression. A two-tailed P-value of less than 0.05 was considered statistically significant. Intra-observer variability was tested by using Spearman’s test and Bland–Altman method.

3. Results

The average age of our TOF patients was 13.1 ± 3.3 (range 7.7–18.7 years). The mean post-operative follow-up time of our patients was 11.8 ± 2.9 years from the corrective surgical procedure (range 7.2–18.4). All patients were of NYHA class I–II and all control patients were of NYHA class I. A total of 16 (40%) of the patients had undergone palliative procedure in the early childhood before the corrective surgery. Transannular patch (TAP) was used in 23 (58%) patients. All the corrective operations were done on cardiopulmonary bypass with aortic crossclamp. The myocardium was protected with cold blood cardioplegia. During the follow-up time a reoperation was done to 6 (15%) of our patients. Table 1 summarizes the baseline characteristics of TOF patients and healthy controls. There were no time-related trends in surgical techniques and all methods were used constantly during the operative period from 1990 to 2003.

Visible RV LGE was present in all of our patients when all segments of RV where studied. Our interest was to explore whether RV LGE could be seen as remote enhancements outside the surgical scars. Therefore, we performed also a LGE score calculation by excluding enhanced areas of the right ventricular outflow tract (RVOT) and the ventricular septal defect (VSD) patch. This remote LGE was found in 39 out of 40 of our patients. Both LV and RV LGE were absent in the control subjects.

Remote LGE was frequently found in all 5 remote segments of RV. Most commonly it was found in the anterior wall (33 of 40 patients) followed by RV insertion points (27/40). In three other segments it
was found as follows: the inferior wall of RV in 16, the RV surface of septum in 15 and in the trabecular bands in 9 patients out of 40.

We found a significant positive correlation between remote RV LGE and RV end-diastolic volume (Fig. 4A). The remote LGE score was significantly higher in patients with severe PR when compared to those without (Fig. 4B). The amount of pulmonary regurgitation also correlated positively with RV EDV ($r = 0.40$, $P = 0.01$).

Since remote LGE was related to pulmonary regurgitation and right ventricular dilation, we studied if post-operative follow-up time would affect LGE. In patients with moderate ($>20\%$) or severe ($\geq 30\%$) pulmonary regurgitation a linear regression analysis demonstrated a correlation between postoperative follow-up time and remote LGE ($r = 0.53$, $P = 0.006$, Fig. 4C). There was a suggestive negative trend between RV EF and LGE score but it did not reach statistical significance.

We then studied whether the levels of two circulating biochemical markers available for diagnostics and follow-up of myocardial disease, namely the NT-proBNP and PIIINP, would parallel RV fibrosis or RV size and function. The levels of NT-proBNP were clearly elevated in our patient group as shown in Table 2. A linear regression analysis demonstrated a correlation between remote LGE score and NT-proBNP levels ($r = 0.38$, $P = 0.015$). In addition, after pooling the data from the control subjects and patients, significant positive correlation existed between NT-proBNP and RV EDV and a negative correlation between NT-proBNP and RV EF (Fig. 5A and B).

Comparison between the patients and control subjects demonstrated that the plasma levels of PIIINP were significantly higher in TOF-patients. However, there was no correlation between PIIINP and CMR-findings or other clinical and biochemical data.

4. Discussion

An important finding of our study was that the RV LGE was present in children with surgically repaired TOF and these LGE changes were more abundant in patients as the right ventricle dilation, age, and follow-up time progressed.

Another new finding of this study was that RV myocardial LGE was found in remote areas of RV of these young patients. We propose that pulmonary regurgitation, significantly present in our patients, and...
LGE has become a marker of fibrotic, scarred or otherwise abnormal myocardium. It has been previously studied in both ischemic and non-ischemic heart disease, and CMR LGE has contributed to understanding LV dysfunction and the pathophysiology behind in these diseases [12,25–27]. Correlation between LGE and myocardial fibrosis has also been validated in earlier histological studies [28,29]. In adult patients with TOF, RV LGE has correlated with adverse clinical markers such as ventricular dysfunction and arrhythmias [13]. Accordingly, parallel to these previously published data we suggest that the adverse hemodynamic loading conditions induced by the post-operative anatomic and hemodynamic conditions are deleterious to the myocardium. In future investigations, it would be interesting to study whether, in the chronic stage of TOF, RV LGE is different in patients with restrictive RV physiology with attenuated RV remodeling [30].

LV LGE was not found in any of our patients. No LV venting was done in any of the patients during the TOF repair. In adult patients with TOF, LGE findings vary: in one study where patients were operated on during adult life, LV LGE was found in 53% of the patients [13]. In another study, no significant LV LGE was encountered in a cohort of 9–67 year old individuals after repair of TOF [15]. Differences in surgical era and the surgical approach, including the omission of left ventricle venting, could explain these differences. Since our patients were children and adolescents, it is also possible that LV LGE develops later in life.

Consistent with previous observations, our present findings demonstrated an increased amount of plasma PIIINP in patients with TOF compared to their age-matched control population. However, PIIINP did not show correlation with LGE, which generally is anticipated to derive at least for some extent from increased fibrosis within the myocardium. In light of the recent literature we feel that this is not surprising since the extracardiac synthesis of fibrillar collagens is significant [31]. In addition, extracardiac turnover of collagens varies not only in noncardiac diseases but also in cardiac disease associated with abnormal loading conditions [32]. These observations may somewhat decrease attractiveness of the circulating collagen-derived peptides as diagnostic and follow-up tools in clinical cardiology [33].

Our study demonstrated a correlation between NT-proBNP and LGE. In parallel with previously published studies in somewhat older patients with TOF [20–22] we found significant correlations between NT-proBNP and RV EDV, pulmonary regurgitation, and RV function. Thus, our present findings add weight on the hypothesis that gradual dilatation of RV after TOF-repair provokes fibrosis within the myocardium, which can be detected by RV LGE.

5. Conclusion

We conclude that RV LGE is present in children after repair of tetralogy of Fallot and it is found also outside the surgically injured areas. The amount of LGE correlates with the severity of PR and RV dilatation. Our data on serum NT-proBNP suggests that RV LGE reflects right
ventricular dilation. We propose that the study of LGE might provide valuable additional information in decision-making for the timing of the operative treatment.

6. Study limitations

Our study was cross-sectional, and a longer follow-up with repeated clinical, radiological and biochemical assessments would be needed to confirm whether the right ventricle fibrosis in children after TOF repair is progressive. Only one marker of fibrous collagen turnover was used.

However, inclusion of type I collagen derived PICP and PINP was not considered due to the potential confounding factors due to the ubiquitous content of type I collagen within the parenchymal organs, skin and the skeleton.

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