Tricuspid three-dimensional ring versus fashioned flexible band annuloplasty in management of functional tricuspid valve regurge: comparative long term study

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Abstract

Background: Functional tricuspid valve regurgitation due to left sided valve lesions remains a common finding. Nowadays, different types of three-dimensional rings and flexible bands are considered to be the gold standard techniques for repair of such functional regurgitation. Our study aimed for long-term comparison of the durability and effectiveness of those techniques.

Results: The data of 170 adult patients with moderate or more functional tricuspid regurgitation secondary to mitral stenosis or regurgitation receiving tricuspid valve repair using either rigid 3D rings (contour 3D or MC3 ring) or prosthetic fashioned bands (Dacron or PTFE) and mitral valve surgery between March 2013 and September 2018 were collected and analysed. Patients were classified in two matched groups: ring group (group A) consisting of 90 cases and band group (group B) consisting of 80 cases. follow-up period ranged from 55.2 to 78 months (mean 67.2 ± 10.8), during this period New York heart association for dyspnea functional class evaluation and full echocardiographic assessment were done and was scheduled as following: before hospital discharge, after 6 months of surgery and finally, every 2 years. In-hospital mortality was 1(1.1%) and 2 (2.5%) with p value 0.49 after ring or band annuloplasty, respectively. No significant differences in both groups regarding hospital mortality and morbidities. There was statistically significant decrease in NYHA class, degree of TR, and PASP within the 2 groups postoperatively with p value of 0.03, 0.04, and 0.033, respectively, with no statistically significant difference between both groups (p value > 0.05). There was statistically significant difference in the freedom from recurrent TR and need for reoperation during the follow-up period between both groups with lower recurrence and reoperation rates in group A, (p value 0.03 and 0.001, respectively).

Conclusion: Tricuspid valve repair with different annuloplasty rigid three dimensional rings or fashioned flexible bands for functional tricuspid regurgitation offers good long-term outcome. Yet, the rigid rings maintain their benefits on the rate of recurrent TR and the need for reoperation.

Keywords: Functional tricuspid regurgitation, Band annuloplasty, Tricuspid 3D ring
Backgrounds
Functional tricuspid regurgitation (FTR) remains a common finding in patients with left-sided valvular heart diseases especially mitral valve regurgitation or stenosis. It is mainly caused by dilatation and flattening of the tricuspid valve (TV) annulus as a result of right ventricular (RV) enlargement and is inevitably affected by the RV function [1]. Multiple studies support the better outcome of using rigid ring annuloplasty over suture repair for treating FTR in concern with the durability and freedom from residual regurgitation. These studies refer the superiority of rigid ring annuloplasty to the three-dimensional configuration achieved by such rings which matches the normal tricuspid annulus and, at the same time, to the rigid support of the annulus [2–6]. On the other hand, other studies support the use of flexible prosthetic bands like Dacron or polytetrafluoroethilin (PTFE), especially over suture techniques, due to their easy applicability and availability with good postoperative results [7, 8]. Therefore, suture anuloplasty became nowadays less convenient for FTR repair since both flexible bands and rigid rings appeared to offer good midterm outcomes in recent meta-analysis studies [9, 10]. However, the ideal annuloplasty method for repairing FTR is still debatable between the three-dimensional (3D) rigid rings versus flexible prosthetic bands regarding the long-term outcome. In this retrospective comparative study, we share our experience with long-term outcome of using 3D rigid rings and flexible fashioned bands in FTR.

Methods
Study method
Between March 2013 and September 2018, a total of 367 patients with moderate or more FTR secondary to mitral stenosis or regurgitation underwent tricuspid valve repair and mitral valve surgery at Cairo University hospitals. Of these, 90 cases received tricuspid valve repair with rigid 3D rings (contour 3D or MC3 ring) while the rest (277 cases) received tricuspid valve repair with prosthetic fashioned bands (Dacron or PTFE). The data of the patients receiving tricuspid valve repair with rigid rings were collected and reviewed (group A). Another group of 80 cases receiving tricuspid valve repair with bands were matched to group A using propensity scores according to preoperative patient characteristics (group B). Cases with organic tricuspid regurgitations, recurrent tricuspid valve disease (TVD), endocarditis affecting the TV, associated tricuspid stenosis, congenital TR, associated aortic or pulmonary valve affection, and concomitant ischemic heart disease were not included in our study. This study was approved by our ethical committee. An informed consent was obtained from all patients for the planned operations and due to its retrospective nature the consent for enrolment of patient data in the study was waived. The indication for surgery for the tricuspid valve in our study was based on the American Heart Association (AHA) guidelines; when TR was severe or if moderate but with tricuspid annular dilatation (‘ 40 mm) or pulmonary artery systolic pressure (PASP) above 50 [11]. The TR severity was assessed by the M mode, two dimension and Doppler echocardiography. Moderate TR (grade 2+) was defined by a width of vena contracta of 4–7 mm and a jet area of 5–10 cm². Severe TR (grade 3+ and 4+) was defined by a width of vena contracta of 7 cm and a jet area of 10 cm² with systolic flow reversal in hepatic veins. Data were categorized into; preoperative, operative, postoperative, and follow-up which ranged from 4.6 to 6.5 years (mean 5.6 ± 0.9). In-hospital mortality was defined by deaths within 30 days of the operation. Recurrent TR was defined by postoperative moderate or more TR (grade 2+ or more).

Data collection:
Full data was recorded and analyzed as following:

Preoperative data: age, sex, New York Heart Association (NYHA) class for dyspnea, transthoracic echocardiographic (TTE) findings including: ejection fraction (EF), left ventricle end diastolic dimensions (LVEDD), left ventricle end systolic dimensions (LVESD), systolic pulmonary pressure (SPAP) in mmHg. We will consider it as mild, moderate, and severe pulmonary hypertension, respectively, with the values of 30–45 mmHg, 45–70 mmHg, and > 70 mmHg.

TR Jet area (cm²), which measures the narrowest distance of the tricuspid jet in mm, tricuspid annular plane systolic excursion (TAPSE), which measures the longitudinal systolic motion of the free edge of the tricuspid valve annulus, tricuspid annulus (TA) diameter, and mid RV diameter in short axis

Operative data
Full records of cardiopulmonary bypass time, cross clamp time, intra-operative findings, type of ring or band in each group, and type of mitral valve intervention.

Postoperative data
Duration of mechanical ventilation, need for pacemaker, intensive care unit (ICU) stay, hospital stay, morbidities, mortality, and follow-up echo data as scheduled by TTE preoperatively as a baseline, then, intra-operative transoesophageal echocardiography (TEE) for correlation with the preoperative echo, postoperative TTE before hospital discharge, then, in outpatient clinic after 6 months of surgery and finally, every 2-year of follow-up
for all patients in both groups with comparison with the preoperative NYHA, TR grade, PASP, and EF. Follow-up period ranged from 55.2 to 78 months (mean 67.2 ± 10.8).

Operative procedure
Surgery was done in all cases through full median sternotomy and cardiopulmonary bypass (CPB) was achieved with routine bicaval cannulation. Myocardial protection was carried out with multi-dose cold blood cardioplegia every 30–40 min delivered in an ante-grade mode with systemic cooling to 32 °C. After aortic cross-clamping the concomitant mitral valve surgery, whether repair or replacement, was done first through left atriotomy. Tricuspid valve repair was then performed through right atriotomy during the rewarming period on flaccid heart with either band (Dacron or Gortex) or using rigid 3D tricuspid rings (Contour 3D [Medtronic, Minneapolis, MN] ring or MC3) which were fixed to the annulus with interrupted non-pledgetted 2/0 non-absorbable (Ethibond) sutures avoiding the area of the AV node. The choice of the type of tricuspid repair was carried out according to the surgeon’s preference. Sizing for rigid rings by different sizers took place according to the base of the septal leaflet (or one size smaller than the anterior tricuspid leaflet surface area). For band annuloplasty, bands of Dacron or Gortex were fashioned in a width of 3–4 mm and a length of 5 to 7 cm depending on the operative assessment of the surgeon. Finally, testing of the repair was done by saline injection into the tricuspid valve while closing the pulmonary valve before the right atrium closure (Fig. 1).

Statistical analysis
A propensity score matching (PSM) analysis to create comparable risk groups for comparing the outcome of using the two different techniques on degree of postoperative tricuspid regurgitation and freedom from reoperation with respect to covariates as demographics and preoperative characteristics. In PSM standardized difference were calculated to compare features before and after matching with imbalance being defined as an absolute value greater than 0.10 (small effect size). Matching was performed using the nearest neighbor algorithm with a calliper distance of 0.0001. All preoperative variables > 90% missing data were excluded from analysis. Data were reviewed and coded to facilitate data manipulation and double entered into Microsoft Access and data analysis was performed using Statistical Package of Social Science (SPSS) software version 25 in Windows 7. Quantitative data were summarized using mean ± standard deviation while categorical data were presented as frequency (count) and relative frequency (percentage). For two independent groups of quantitative parametric data non-paired Student’s t test was used while for qualitative data chi-squared test was used. Two-way ANOVA test was implied for repeated measures. A p value < 0.05 was considered the cut-off value for significance.

Results

Early results
Preoperative patient characteristics are summarized in Table 1 including epidemiological, clinical, and echocardiological data. There was no statistical significant difference between both groups, which indicates proper

| Table 1 Preoperative data in both study groups |
| Variables | Ring group | Band group | p value |
| --- | --- | --- | --- |
| Sex | | | |
| Male | 33(36.7%) | 28(35%) | 0.8 |
| Female | 57(63.3%) | 52 (65%) | |
| Age (years) | 39.7 ± 6.5 | 37.9 ± 7.5 | 0.08 |
| TR grads | | | |
| II | 36(40%) | 24(30%) | 0.15 |
| III | 18(20%) | 12(15%) | |
| IV | 36(40%) | 44(55%) | |
| Mitral valve lesions | | | |
| Stenosis | 74 (82%) | 65 (81%) | 0.75 |
| Regurgitation | 4 (5%) | 2 (2.5%) | |
| Stenosis and regurgitation | 12 (13%) | 13 (16.5%) | |
| NYHA class | 2.9 ± 0.85 | 3.1 ± 0.92 | 0.14 |
| LVEF % | 53.3 ± 16.8 | 52.0 ± 18.9 | 0.49 |
| LVSD (cm) | 3.7 ± 0.98 | 3.92 ± 0.53 | 0.07 |
| LVEDD (cm) | 6.04 ± 0.98 | 5.87 ± 0.56 | 0.16 |
| TAPSE (cm) | 1.8 ± 0.43 | 1.9 ± 0.3 | 0.08 |
| RV short axis (cm) | 2.93 ± 0.37 | 2.99 ± 0.4 | 0.13 |
| TA diameter (cm) | 4.8 ± 0.4 | 4.7 ± 0.5 | 0.15 |
| PASP (mmHg) | 63.52 ± 9.95 | 61.23 ± 10.25 | 0.14 |

TR tricuspid regurgitation, NYHA New York Heart Association, LVEF left ventricular ejection fraction, LVEDD left ventricle end diastolic dimensions, LVESD left ventricle end systolic diameter, TAPSE tricuspid annular plane systolic excursion, RV right ventricular, TA tricuspid annulus, SPAP systolic pulmonary artery pressure
matching between the two study groups. The pathology of the mitral valve affection was rheumatic in origin in all cases of the study. The sizes of the rigid rings (MC3 in 20 cases and Contour 3D in 70 cases) used in group A ranged from 28 to 34 cm with a mean value of 30.5 cm. On the other hand, fixed lengths of 5 to 7 cm of flexible fashioned bands (Dacron in 70 cases and Gortex in 10 cases) were used in group B. There was statistical significant difference of bypass time and cross clamp time being shorter in group B. Other operative and postoperative parameters are shown in Table 2 with no statistically significant differences between both groups. One case (1.1%) died in group A due to multi-organ failure while two cases (2.5%) died in group B from refractory low cardiac output in one patient and respiratory failure due to pneumonia and prolonged ventilation in the other case with no significant difference. On comparison between the preoperative and postoperative values for NYHA class, degree of TR, and PASP, there was statistically significant decrease in the immediate postoperative period within the 2 groups with a p value of 0.03, 0.04, and 0.033, respectively. However, there was no statistically significant difference between both groups (p value = 0.065).

Late results and follow-up
Our complete long-term follow-up was achieved in 134 (78.82%) cases due to mortalities and patients non-compliance (8.8%). Follow-up period ranged from 55.2 to 78 months (mean 67.2 ± 10.8). Late deaths occurred in 18 (10.58%) of cases. Ten cases in group A as well as in 8 case in group B due to infective endocarditis in 4 cases, stuck mitral valves in 3 cases, cerebrovascular events in 3 cases, renal failure in 2 cases, malignancy in 2 cases, and 4 unknown causes with no statistically significant difference (Fig. 2).

There was no statistically significant difference in the mean NYHA class during the follow-up period between both groups (0.91 in group A and 0.92 in group B) with a p value of 0.09 (Fig. 3). Also, PASP and EF% showed no statistical difference in the same follow-up periods between both groups. However, there was a statistically significant difference in the mean degree of TR improvement at the same follow-up period in favor of group A (1 in group A and 1.5 in group B) with a p value of 0.03. The degree of improvement and severity of TR in the follow-up period is shown in Fig. 4.

In addition, 7 cases (8.75%) out of the 80 patients in group B and 4 (4.4%) out of the 90 patients in group A had recurrent TR during follow-up period. Four patients from the 7 cases with recurrent TR in the band group needed reoperation with replacement of the tricuspid valve due to recurrent right sided heart failure despite maximal medical treatment. On the other side, the four patient with recurrent TR in ring group continued their follow-up period without reoperation. There was statistically significant difference in the freedom from recurrent TR and need for reoperation during the follow up period.

| Table 2 | Operative and outcome data in both study groups |
|---------|------------------------------------------------|
| Variables | Ring group (n = 90) | Band group (n = 80) | p value |
| Bypass time (min) | 116 ± 25.41 | 105.6 ± 20.33 | < 0.001* |
| Cross clamp time (min) | 91 ± 28.6 | 76.7 ± 30.2 | < 0.001* |
| Concomitant mitral valve surgery | Repair: 6(6.7%) | 4(5%) | 0.64 |
| Replacement | 84(93.3%) | 76(95%) | |
| Types of ring or band | MC3 20: (22.2%) | Dacron: 70(87.5%) | |
| Contour 3D: 70 (77.8%) | Gortex: 10(12.5%) | |
| Postoperative mechanical ventilation (h) | 8.1 ± 1.9 | 7.7 ± 1 | 0.08 |
| ICU stay (days) | 2.9 ± 0.76 | 2.6 ± 1.3 | 0.07 |
| Hospital stay (days) | 7.4 ± 1.2 | 7.8 ± 1.5 | 0.06 |
| Postoperative complications | Reoperation for bleeding | | 0.9 |
| Renal failure | 2(2.2%) | 2(2.5%) | |
| Cerebrovascular stroke | 1(1.1%) | 1(1.3%) | |
| New pacemaker insertion | 1(1.1%) | 0(0%) | |
| Tracheostomy | 0(0%) | 1(1.25%) | |
| Pneumonia | 0(0%) | 2(2.5%) | |
| In-hospital mortality | 1(1.1%) | 2(2.5%) | 0.49 |
| Reoperation for recurrence | 0(0%) | 4(5%) | 0.001* |

ICU intensive care unit
between both groups with lower recurrence and reoperation rates in group A with a $p$ value of 0.03 and 0.001 respectively (Fig. 5).

Discussion
Secondary FTR is very common at the time of left sided valve surgery due to progressive annular dilatation mainly leading to lack of proper tricuspid leaflets coaptation and regurge. Furthermore, the changes that occurs in valve geometry and the more planer the tricuspid annulus that become with prolonged dilatation.

Current data suggest that the dilatation that occurs in cases of FTR especially with left-sided valve lesions is an on-going process that continues to progress unless surgical annuloplasty is done at the time of left-sided valve surgery to prevent future progression of the disease [12]. Without intervention, persistent TR is currently associated with poor outcome even if not sever and therefore, the recommendations for repairing the FTR nowadays changed to lesser threshold for regurgitation [13].

The main difference between both rigid 3D rings and flexible fashioned bands is that rigid rings restore the saddle, non-planner, and tricuspid annulus which have been proven by the studies using 3D echocardiography to improve annular symmetrical contractions during the cardiac cycle. In addition, rigid rings seem to give more long-term stability of the tricuspid annulus [14]. On the other hand, flexible bands maintain the physiological motion of the tricuspid annulus during cardiac contractions which is now well-recognized by many researches but its clinical importance and role are not yet fully investigated [15].

In our study, the mean size of used rigid rings was 30.5 (range 28–34). We simply used one size smaller of the 3D ring (contour 3D or MC3) since some studies showed that the exact annular size is relatively smaller when the heart is beating than when arrested. Also,
other studies recommended routine downsizing in all cases whether performing the tricuspid repair on beating or arrested heart with comparable results [3, 16]. However, multiple studies failed to identify a concrete relation between the ring size, as a risk factor, and recurrent TR with no much benefits of aggressive downsizing [17, 18].

On the other hand, we fashioned the bands to a width of 3 mm and a length of 5 to 7 cm to plicate the anterior and posterior annulus depending upon the intraoperative measurement of the tricuspid annulus and the degree of TR. Although the fact that there are no solid guidelines for this measurement, but this length has been obtained in previous studies from the average circumferences of most used ring sizers. In addition, it has been shown in previous studies that the tricuspid annulus measured by preoperative echocardiography from the mid-anterior to the mid-septal leaflet in the end-diastole using the apical four chamber view differs from the transvers annulus measured surgically from the antero-septal commissure to the antero-posterior commissure. For example, an annulus estimated by preoperative echocardiography to be 4 cm would be measured surgically 7 cm [19].

In the band group, we used Dacron or Goretx assuming that both has nearly equal flexibility with fashioning the band 3 mm width and from 5 to 7 cm length depending upon the intraoperative measurement of the tricuspid annulus and to what degree the TR present and the needs for the proper downsizing with no solid guidelines for measurement but, we took this measurement previously from average circumferences of most used ring sizers.
In our series, we have succeeded effectively to reduce the tricuspid annulus size by either the rigid rings or fashioned flexible bands with no statistically significant difference in both modalities in the aspect of improving the TR degree, PASP and NYHA class in the early post-operative period. However, rigid 3D rings significantly maintained long-term effects on the repair with less re-intervention and recurrent TR during our follow-up. This finding agrees with previous multiple studies and meta-analysis comparing rigid rings with flexible band devices, not fashioned (custom made) bands, as we tried. We use these intra-operatively fashioned bands for easy accessibility and applicability with similar concepts [9, 10, 20–22]. Moreover, some studies used native pericardium fashioned into similar bands as it is widely available for all patients with encouraging results [8, 23].

Nevertheless, most of the studies addressing the risk factors for recurrent TR after surgical repair of FTR reported LV dysfunction, high preoperative PASP and the presence of tricuspid valve tethering [24, 25] these studies could not identify the type of annuloplasty as a risk factor for recurrent TR which implies the importance of multi-centric prospective studies with long term results. We could not include TV tethering in our preoperative echo parameters, and we did not study the risk factors for recurrence as PASP, also, mean LVEF was in normal range in our preoperative data and maintained in the long-term follow-up with no significant changes to be a risk factor.

Study limitations
Our study has several limitations. The retrospective nature of the study and the sample size was relatively small. There is procedural difference between fashioned bands which cover the anterior and posterior annulus only and rigid rings which cover in addition part of septal annulus. There were no standard guidelines for sizing the fashioned tricuspid bands. The risk factors for recurrence have not been studied. Finally, the study did not compare different annuloplasty material used in bands or different manufactured rigid rings.

Conclusion
Tricuspid valve repair with different annuloplasty rigid three-dimensional rings or fashioned flexible bands for functional tricuspid regurgitation offers good long-term outcome. Yet, the rigid rings on the long run maintain their benefits on decreasing the rate of recurrent TR and the need for reoperation.

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Authors' contributions
All authors participated in getting the conception, data collection, analysis, and manuscripts writing. ME was a major contributor in analysis and interpreted the patient data regarding the comparative parameters. AN was a major contributor in writing the manuscript. SN was major contributor in revising and editing the manuscript. All authors read and approved the final manuscript.

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Consent for publication
Due to the retrospective nature patients, consent for publication was waved.

Competing interests
The authors declare that they have no competing interests.

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Abbreviations
AHA: American heart association; CPB: Cardiopulmonary bypass; EF: Ejection fraction; FTR: Functional tricuspid regurgitation; ICU: Intensive care unit; LVEDD: Left ventricle end diastolic dimensions; LVESD: Left ventricle end systolic dimensions; NYHA: New York Heart Association; PTFE: Polytetrafluoroethilin; RV: Right ventricular; SPAP: Systolic pulmonary pressure; SPSS: Statistical Package of Social Science; TR: Tricuspid regurgitation; TDO: Tricuspid valve disease; TTE: Transthoracic echo; TAPSE: Tricuspid annular plane systolic excursion; TA: Tricuspid annulus; TEE: Trans-oesophageal echocardiography; 3D: Three dimensions

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