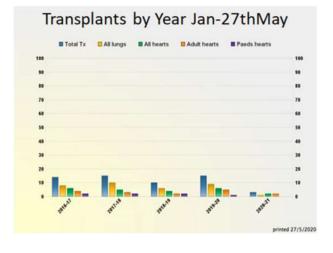


Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active. transplant activity for each financial year up to April 2020 and our activity between Jan - 27th May 2020 (the first UK pandemic peak) comparing this to the same period over the last five years. This allowed us to directly compare the effect of COVID with previous years activity.

Results: A total of 19 heart patients died during Jan to 27th May 2020. Out of the total deaths 4 were from paediatric group, 11 were post-transplant and 5 on the active list. There were no cardiac deaths related to COVID 19. The total deaths during the same period were 19, 24, 20 and 32 in the years 2019, 2018, 2017 and 2016 respectively. A total of 41 lung patients died during Jan to 27th May 2020. Out of the total 11 were on the active list awaiting transplant and 19 were post-transplant deaths. Of these 2 were COVID deaths. The corresponding deaths during year 2019, 2018, 2017 and 2016 were 46, 51, 49 and 46 respectively. Transplant activity from Jan to 27th May in 2016-19 averaged 32 (16 hearts,16 lungs); activity during the same time period in 2020 was 4 lungs and 13 hearts, clearly affected by reduced activity due to COVID19.

Conclusion: This audit does not suggest a significant rise in mortality due to the COVID 19 pandemic in vulnerable patients either pre or post-transplantation. Measures such as shielding were highly effective in this population. Transplant activity for the year was affected.



(26)

Rapid Virtualization of a Heart Function Clinic in Response to the COVID-19 Pandemic

M. Linghorne, M. O'Sullivan, Y. Moayedi, N. Aleksova, D. Delgado, A.C. Luk, M.A. McDonald, F. Billia, H.J. Ross and J.G. Duero

Posada. Cardiology, University Health Network, Toronto, ON, Canada.

Purpose: The current COVID-19 pandemic has had an unprecedented impact on healthcare systems across the world. It has stretched to the limit acute care systems, indirectly it has shaped new and innovative ways to deliver care for those with chronic conditions. Herein we describe initial outcomes of the rapid virtualization of the Heart Function Clinic at a major quaternary Hospital in Toronto, Ontario.

Methods: Consecutive patients attending the heart function clinic at the Toronto General Hospital between March 9, 2020 and June 30, 2020 were included. Visits were classified as "in-person" if patients were physically present for the clinical interaction and "virtual" if the clinical interaction occurred while the patient was away using currently available modes of communication: telephone or web-enabled (Ontario Telemedicine Network -OTN, or other available web-based applications). The purpose of the individual visit was categorized as: "surveillance", "titration", "new assessment" or "Clinical trial".

Results: A total of 292 patients had a total of 521 clinical encounters during the lockdown period. Of these, 168 (32.2%) were "in-person", while 353 (67.8%) were "virtual". 101 (19.3%) were primarily for the purposes of titration. These virtual assessments led to 14(2.7%) in-person assessments. 258 (49.5%) of patients had an LVEF < 40%, among these patients

220 (85.3%) were on an ACEi, ARB or ARNi, 242 (93.8%) on a Betablocker, 191 (74%) on an MRA, 46 (17.8%) on SGLT2inhibitor.

Conclusion: Rapid virtualization of a large academic multi-disciplinary clinic is possible. This allows for ongoing delivery of safe care to patients with chronic conditions and can be used as a model for other clinics facing the pandemic. Lessons learned will be used to transition to a hybrid model of in-person and virtual even after the pandemic has come to an end.

(27)

Telemedicine (TM) during SARS-CoV-2 Outbreak

<u>M. Masetti,</u> S. Toniolo, A. Adorno, L. Giovannini, P. Prestinenzi, M. Sabatino, A. Russo, S. Martin Suarez, A. Loforte, D. Pacini and L. Potena. Bologna Academic Hospital, Bologna, Italy.

Purpose: As Italy faced SARS-CoV-2 outbreak as first country outside China, and our hospital converted most of activities into the ones for COVID-19 patients (pts), we had to manage the need for continuing care of advanced heart failure (HF), heart transplant (HT) and LVAD pts. TM was a possible strategy, but its role in this very sick cohort is unknown.

Methods: During the lockdown (03-05/2020), we decided to make either a phone (PV) or an in presence (IV) visit, selecting for IV pts listed for HT, with LVAD, recently HT, scheduled for a biopsy within 6 months after HT or a RHC for listing eligibility. In PV, we assessed symptoms, blood pressure, drugs, and programmed a subsequent IV. All pts in IV group were triaged by phone for COVID-19 symptoms or contacts and if scheduled for RHC or biopsy received SARS-CoV-2 swab 48 h before the procedure. Study endpoints were: combined incidence at 6 months of MACE (HF hospitalization, CV death and need for anticipated IV) in HF/VAD group, and MACE, rejection and any cause- hospitalization in HT group.

Results: Among 448 pts (57 \pm 12y, 240 HT, 191 HF, 17 LVAD), 52% were managed by PV and a subsequent IV was scheduled after 3 \pm 2 months. Pts managed by PV were healthier: in HF-VAD group they were less frequently listed, had less Afib, LVAD (2/17) (p<0.01 all); post-capillary PH (pC-PH) was similarly distributed; in HT group there were less pts transplanted in the last 5 years (15% vs 52%, p<0.01) and numerically less with 2R rejection in the previous 6 months (8.3% vs 27.1%, p=0.13).The PV group had a lower incidence of the endpoints in both HF/VAD and HT cohorts (92.3 \pm 2.3% vs 70.3 \pm 4.4%; 97.0 \pm 1.7%vs82.5 \pm 4.1%, p<0.01). Overall, the predictors of the endpoints at multivariate analysis were pC-PH and PV (HR: 5.2 and 0.1, p<0.03 both) and a recent 2R rejection (HR: 3.6, p=0.05) in the HF/VAD and HT group respectively.There were no cases of COVID-19 in IV; 5 pts got infected at home in a context of infection prevalence of 6/1000 inhabitants in our region and of 40% of hospital beds dedicated to COVID-19 pts.

Conclusion: In this retrospective study, by reporting an organization set up in a emergency situation, we show that TM can be safely used to manage stable HF, LVAD and HT patients, whereas pC-PH and a recent rejection may identify those needing IV. These data suggest that the availability of devices for monitoring pulmonary pressures may improve safety of PV in HF pts and that TM could be useful not only in a pandemic outbreak but also subsequently.

(28)

The Effect of Body Mass Index on Presentation of COVID-19 amongst Heart Transplant Recipients: A Multi-Institutional Study

<u>A. Iyengar,</u>¹ J.J. Han,¹ M.R. Helmers,¹ B.F. Smood,¹ W.L. Patrick,¹ J. Kelly,¹ N. Moss,² S.S. Najjar,³ B.A. Houston,⁴ R.J. Tedford,⁴ S. Shore,⁵ E. Vorovich,⁶ E. Hsich,⁷ K.M. Alexander,⁸ S. Chaudhry,⁹ H. Vidula,¹⁰ A. Kilic,¹¹ M.V. Genuardi,¹² E.Y. Birati,¹ and P. Atluri.¹ ¹Penn Medicine, Philadelphia, PA,² The Zena and Michael A. Wiener Cardiovascular Institute, Icahn School of Medicine at Mount Sinai, New York, NY; ³MedStar Washington Hospital Center, Washington, DC; ⁴Medical University of South Carolina, Charleston, SC; ⁵Cardiovascular Division, University of Michigan, Ann Arbor, MI; ⁶Division of Cardiology, Northwestern University, Chicago, IL; ⁷Heart and Vascular Institute at the Cleveland Clinic, Cleveland Clinic Lerner College of Medicine of Case Western Reserve University SoM, Cleveland, OH; ⁸Division of Cardiovascular Medicine and the Stanford Cardiovascular Institute, Stanford University School of Medicine, Stanford, CA; ⁹St. Vincent Medical