10th September 2019
Staff Publications
To request an article, please email health.library@sa.gov.au
Links to abstract where available.

Department for Health and Wellbeing

Chew DP; Lambrakis K; Blyth A; Seshadri A; Edmonds MJR; Briffa T; Cullen LA; Quinn S; Karnon J; Chuang A; Nelson AJ; Wright D; Horsfall M; Morton E; French JK; Papendick C;
A Randomized Trial of a 1-Hour Troponin T Protocol in Suspected Acute Coronary Syndromes: The Rapid Assessment of Possible ACS In the Emergency Department with High Sensitivity Troponin T (RAPID-TnT) Study.
Circulation. 2019 Sep 03.

Royal Adelaide Hospital

Singh JP; Abraham WT; Auricchio A; Delnoy PP; Gold M; Reddy VY; Sanders P; Lindenfeld J; Rinaldi CA
Design and rationale for the Stimulation Of the Left Ventricular Endocardium for Cardiac Resynchronization Therapy in non-responders and previously untreated patients (SOLVE-CRT) trial.

Elliott AD; Mishima RS; Lau DH; Sanders P
Improving Exercise Tolerance with Catheter Ablation

Pham H; Trahair L; Phillips L; Rayner C; Horowitz M; Jones K
A randomized, crossover study of the acute effects of acarbose and gastric distension, alone and combined, on postprandial blood pressure in healthy older adults.

Chan MM; Tapia Rico G
The “pet effect” in cancer patients: Risks and benefits of human-pet interaction. [Review]

Mittal P; Price ZK; Lokman NA; Ricciardelli C; Oehler MK; Klingler-Hoffmann M; Hoffmann P
Matrix Assisted Laser Desorption/Ionization Mass Spectrometry Imaging (MALDI MSI) for Monitoring of Drug Response in Primary Cancer Spheroids.

De Sousa SMC; Wang PPS; Santoreneos S; Shen A; Yates CJ; Babic M; Eshraghi L; Feng J; Koszyca B; Roberts-Thomson S; Schreiber AW; Torpy DJ; Scott HS
The Genomic Landscape of Sporadic Prolactinomas.
Royal Adelaide Hospital cont.

Peukert D; Kempson I; Douglass M; Bezak E
Gold Nanoparticle Enhanced Proton Therapy: Monte Carlo Modeling of Reactive Species' Distributions Around a Gold Nanoparticle and the Effects of Nanoparticle Proximity and Clustering. 

Gallagher C; Hendriks JM; Middeldorp ME; Elliott AD; Lau DH; Sanders P
Reducing the Burden of Atrial Fibrillation Cost: Is Integrated Care the Answer? 

Munoz MA; Jurczylik J; Simon A; Hissaria P; Arts RJW; Coman D; Boros C; Mehr S; Rogers MJ
Defective Protein Prenylation in a Spectrum of Patients With Mevalonate Kinase Deficiency. 

Ryan TG; Curragh DS; Ellis D; Selva D; Davis G
Orbital sarcoidosis with bony destruction. 
*Clinical & Experimental Ophthalmology*. 2019 Sep 06.

Haude M; Lee SWL; Worthley SG; Silber S; Verhey S; Rosli MA; Botelho R; Sim KH; Abizaid A; Mehran R; REMEDEE Trial Investigators
The REMEDEE trial: 5-Year results on a novel combined sirolimus-eluting and endothelial progenitor cells capturing stent. 
*Catheterization & Cardiovascular Interventions*. 2019 Sep 05.

Cockbain AJ; Parameswaran R; Watson DI; Bright T; Thompson SK
Flatulence After Anti-reflux Treatment (FAART) Study. 

SA Health

Banham D; Roder D; Keefe D; Farshid G; Eckert M; Howard N; Canuto K; Brown A; CanDAD Aboriginal Community Reference Group and other CanDAD investigators
Disparities in breast screening, stage at diagnosis, cancer treatment and the subsequent risk of cancer death: a retrospective, matched cohort of aboriginal and non-aboriginal women with breast cancer 
Motum P; Just S; Zebeljan D; Nicholls C; Kershaw G; Oliver S; Mohammed S; Favaloro EJ

A diagnosis of von Willebrand disease despite normal test results for factor VIII and von Willebrand factor antigen and activity.


Mucha BE; Banka S; Ajeawung NF; Molidperee S; Chen GG; Koenig MK; Adejumo RB; Till M; Harbold M; Perrier R; Lemyre E; Boucher RM; Skotko BG; Waxler JL; Thomas MA; Hodge JC; Gecz J; Nicholl J; McGregor L; Linden T; Sisodiya SM; Sanlaville D; Cheung SW; Ernst C; Campeau PM

A new microdeletion syndrome involving TBC1D24, ATP6V0C, and PDPK1 causes epilepsy, microcephaly, and developmental delay.

Genetics in Medicine. 21(5):1058-1064, 2019 05.

Magistroni V; Mauri M; D’Aliberti D; Mezzatesta C; Crespiatico I; Nava M; Fontana D; Sharma N; Parker W; Schreiber A; Yeung D; Pirola A; Readelli S; Massimino L; Wang P; Khandelwal P; Citterio S; Viltadi M; Bombelli S; Rigolio R; Perego R; Boulwood J; Morotti A; Saglio G; Kim DW; Branford S; Gambacorti-Passerini C; Piazza R

De novo UBE2A mutations are recurrently acquired during chronic myeloid leukemia progression and interfere with myeloid differentiation pathways.

Haematologica. 104(9):1789-1797, 2019 Sep.

Mucha BE; Banka S; Ajeawung NF; Molidperee S; Chen GG; Koenig MK; Adejumo RB; Till M; Harbold M; Perrier R; Lemyre E; Boucher RM; Skotko BG; Waxler JL; Thomas MA; Hodge JC; Gecz J; Nicholl J; McGregor L; Linden T; Sisodiya SM; Sanlaville D; Cheung SW; Ernst C; Campeau PM

Correction: A new microdeletion syndrome involving TBC1D24, ATP6V0C, and PDPK1 causes epilepsy, microcephaly, and developmental delay.