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台灣移植醫學學會教育訓練課程

2021/12/11(六) **②** 高雄展覽館3樓 304B會議室

Time	Topic	Speaker	Moderator
13:30-13:45	報到		
13:45-14:15	[Polyomavirus BK infection in immunosuppressed patients] Sceening and Diagnosis of polyomavirus BK infection in kidney transplantation	陳呈旭 ^{台中榮總}	張浤榮
14:15-14:45	[Polyomavirus BK infection in immunosuppressed patients] Management of polyomavirus BK-associated nephropathy	田亞中 林口長庚	賴彬卿
14:45-15:00	Coffee Break		
15:00-16:00	【COVID專題演講】 COVID-19 and Kidney Transplantation	張勝勛 ^{成大醫院}	吳麥斯
16:00-17:00	【COVID專題演講】 Managing immunosuppressants in liver transplantation – what have we learnt from COVID-19 pandemic?	陳登偉 ^{三軍總醫院}	胡瑞恒
17:00-18:00	第十三屆第五次理監事會議	王 叙涵 ^{林口長庚}	吳麥斯

台灣移植醫學學會 2021 移植年會



Time	Topic	Speaker	Moderator
08:30-09:00	報到		
09:00-10:00	【特別演講】 Optimizing Immunosuppression: To prevent DSA/AMR and BKVN in Kidney Transplantation	Prof. Peter Nickerson University of Manitoba ON	王叙涵
10:00-10:30	【特別演講】 Sedative and Immunosuppressive Effects of Dexmedetomidine in Transplantation	李正方 ^{林口長庚}	李威震
10:30-11:00	Coffee Break		
11:00-11:30	【Kidney專題】 Long-term outcomes of living kidney donors: lessons learned from Taiwanese nationwide cohort	吳明儒 台中榮總	江仰仁
11:30-12:00	【Liver專題01】 Peri-transplant treatment strategies for HCC beyong UCSF criteria	李明哲 ^{萬芳醫院}	石宜銘 謝宗保
12:00-13:00	[Lunch Symposium] Challenge the Past, A New Clinical Practice, A New Standard of care	Prof. Federico Oppenheimer Hospital University of Barcelona	吳明儒
13:00-13:30	會員大會	王叙涵 ^{林口長庚}	吳麥斯
13:30-14:00	【Liver專題02】 Liver transplantation for hepatocellular carcinoma: concern and perspective	詹昆明 ^{林口長庚}	何明志 林毅志
14:00-14:30	Coffee Break		
14:30-15:00	【Heart專題】 Organ care system in heart transplantation	林宜璋 ^{三軍總醫院}	蔡建松 羅傳堯
15:00-16:00	【特別演講】 ABOi Liver transplant management in AMC	Prof.Pgi-Won Song Asan Medical Center	李威震

台灣移植醫學學會 2021 移植年會

🔛 2021/12/12(日) 👤 高雄展覽館3樓 306會議室

Time	Topic	Speaker	Moderator
08:30-09:15	報到		
09:15-09:30	心臟移植者接受阿斯利康COVID-19疫苗後 產生移植後淋巴組織增生性疾病個案報告	湯文睿 成大醫院	
09:30-09:45	腸引流的胰臟移植:台北榮民總醫院之經驗	施沐姍 ^{台北榮總}	n+ ++
09:45-10:00	器官捐贈之性別差異	廖麗鳳 台北榮總	陳芸 李芳艷
10:00-10:15	Severe herpes zoster infection in patients with solid organ transplantation: a nationwide population-based cohort study with propensity score matching analysis	游棟閔 ^{台中榮總}	王水深
10:15-10:30	病友手冊與個案管理電子化之整合性平台	林姿妤 ^{台北榮總}	
10:30-11:00	Coffee Break		
11:00-11:15	利用大隱靜脈植體之活體親屬腎臟移植:個案報告	陳韋辰 ^{台中榮總}	
11:15-11:30	小兒雙側腎臟挶贈	楊涵中 ^{台中榮總}	闕士傑
11:30-11:45	單一醫學中心高齡活體捐贈者腎臟移植之安全性與受贈者預後經驗	郭芳成 ^{台北榮總}	余家政
11:45-12:00	單一醫學中心使用腦死小兒捐贈者腎臟移植之經驗	廖麗鳳	
12:00-13:00	【台灣腎臟醫學會】Lunch Symposium		
13:00-13:30	會員大會 304B會議室		
13:30-13:45	運用肝臟移植治療晚期肝癌的曙光乍現: 以免疫療法做為移植前降期治療的策略	李明哲 ^{萬芳醫院}	
13:45-14:00	單一醫學中心在活體肝臟移植中使用雙右肝門脈右肝植體之經驗	陳正彥 ^{台北榮總}	李威震 陳登偉
14:00-14:15	右肝活肝移植使用人工血管重建靜脈的術後併發症: 單一醫學中心病例系列報告	黃詩雲 ^{彰基醫院}	171.32.14
14:15-14:30	Coffee Break		
14:30-14:45	腎移植患者感染新冠病毒後的死亡率, 急性腎損傷及失去移植腎功能:縱論與統合分析研究	吳欣旭 ^{林口長庚}	
14:45-15:00	心死捐贈之腎臟移植結果報告:台中榮總經驗分享	張家程	
15:00-15:15	C肝捐贈者心臟停止後死亡的腎臟捐贈	李宗穎	尹文耀
15:15-15:30	多瘤病毒腎病變患者泌尿道腫瘤的高發生率及早期發生的現象	田亞中	張勝勛
15:30-15:45	腎移植病人BK病毒感染與腎病變的案例系列報告分享	詹秀珍 ^{雙和醫院}	
15:45-16:00	Monoclonal antibody use in kidney transplantation patient with Critical COVID 19 infection	游博翰 ^{雙和醫院}	





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陳呈旭 Chen ,Cheng-Hsu

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Diagnosis of polyomavirus infection in kidney transplantation 賢移植中多瘤病毒感染的診斷

多腫瘤病毒BK病毒於1971年首次分離出來一直未被重視,直到是腎移植功能障礙和 腎移植功能喪失的重要危險因素。自1995年匹茲堡大學Randhawa第一次發表多腫瘤 病毒(BK病毒)造成腎臟腎炎才開始受到大家的注意。而腎移植受贈者BKVAN發生 率約為2%至8%,經常導致移植腎功能喪失的重要原因。幾乎大多數情況下,BKV是 多瘤病毒腎病的主因,但還是有少數病人可能檢測到是JC病毒造成,致病機轉和致 病力尚需探討。到目前為止BK病毒感染的治療選擇仍然十分有限,並且沒有有效的 預防措施。儘管過度免疫抑制仍然是移植後BK感染的主要危險因素,但男性、受贈 者年齡較大、先前的排斥反應、人類白細胞抗原錯配程度、冷缺血時間延長、BK 血 清狀態和輸尿管支架置入都被認為是危險因素。BKV感染可能造成原因或許原發感染 或由捐贈者帶入體內造成雙重感染(捐贈者傳播)或BKV再活化感染造成,因為大多 數成年人是BKV血清陽性反應。而異體腎移植排斥,反覆的抗排斥藥物治療也是可能 的一個重要的誘發因素。多瘤病毒腎病患者通常沒有發燒或其他感染症狀,僅表現 為血清肌酸酐升高。BK的常規篩查已被證明可有效預防BK病毒尿或病毒血症病人的 異體腎移植物功能喪失的方法。減少免疫抑制依然是BK腎病治療的主要方法,也是 研究得最好的介入措施。在無法執行BKV篩檢時,在尿液細胞學檢查出decov cell常是 一個重要線索,BKV活化後會在腎小管和尿路上皮細胞的細胞核內增殖,導致細胞核 變大及出現核內包涵體,此包涵體內會有均質化的嗜鹼性顆粒存在,且包涵體外圍 會有染色質沉積在周圍的現象,此種細胞稱為Decoy cell;以尿液細胞學中的 Decoy cell 作為檢測 BKV 感染和預警 BKVAN 在各項研究已經證實有很好的敏感度及不錯的 特異性,是一項簡單敏感的檢驗方法;但尿液中的 Decov cell仍是屬於多變的型態, 容易被誤認為異常或是惡性細胞。藉由血液中BKV的PCR診斷BKV或JCV及其定量PCR (quantitative PCR: qPCR)進行定量來強化診斷,且病毒載量越高,病毒相關之腎臟病 的特異性就越高。然而,腎臟切片檢查仍是目前的標準診斷,可以明確診斷、疾病 分期(對預後有影響),並用於鑑別可能相伴隨疾病(如急性排斥)過程。多瘤病毒在 賢移植病人之發病率、危險因素、臨床相關聯因子和其預後,仍是一個移植賢功能 喪失很值得重視和探討的的議題。



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2018王民寧基金會學術審查委員

多瘤病毒腎病變的處置

An epidemiologic study demonstrated that 80-100% of the general population was infected by BK polyomavirus (BKV) in their childhood. Following primary infection, BKV is persistently hidden in the renourinary tract, and when the immunity of the organ transplant recipients is suppressed, BKV rapidly replicates to cause renal inflammation, the occurrence of BK viruria and BK viremia, and sometimes BKV-associated nephropathy (BKVAN). Potent immunosuppressants reduce acute rejection episodes but increase the risk of BKVAN and its associated allograft loss. A reduction or modification of immunosuppressants may alleviate the progression of BKVAN. Recently, our study and other studies demonstrated a high incidence of urinary tract cancers in BKVAN patients, suggesting an association between BKV infection and urinary tract cancer development. Therefore, early intervention and management of BKV infection are critical for prevention of BKVAN development and subsequently allograft loss. In this lecture, the possible risk factors and triggers for BKVAN development will be discussed. In addition, targeting on these potential factors that may alleviate BKV infection will be introduced. Finally, the effectiveness of different strategies including modification or reduction of immunotherapy in management of BKVAN will be compared.



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移植免疫; 腎臟移植; 腎臟缺血再灌流損傷

COVID-19 and Kidney Transplantation

The COVID-19 pandemic has a great influence on organ donation and transplantation. This viral infection impacts more severely kidney transplant recipients than general population. COVID-19 vaccination remains the primary health strategy to prevent SARS-CoV-2 infection. The clinical trials that evaluated the safety and efficacy of the COVID-19 vaccines excluded solid organ transplant recipients. According to published data, seroconversion after mRNA SARS-CoV-2 vaccination might be unsatisfactory in kidney transplant recipients. The SARS-CoV-2 vaccine efficacy in this subgroup patients is reduced. The impaired humoral and cellular immunity may be caused by immunosuppression. Discordance between humoral and cellular response were also found, however, it is unknown if seronegative patients develop at least a cellular response that could offer a certain grade of protection against SARS-CoV-2. The severity of COVID-19 could potentially be affected by the type and the intensity of the immunosuppressive drugs. The management of immunosuppressive therapy in kidney transplant recipients affected by SARS-CoV-2 may require an individualized approach.

Adjustments to the immunosuppressive regimen are necessarily based upon disease severity, time posttransplant, and the risk of acute allograft rejection. COVID-19 poses challenges for kidney transplantation, this section reviews some issues related to kidney transplantation and aspects of COVID-19 care.



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1999 – 2000 Attending staff surgeon, Division of General Surgery, Tri-Service General Hospital

2000- 2001 Visiting scholar in Clinical Multi-organ Transplantation, The Dumont-UCLA Transplant Center

2003-2004 Director of Gastrointestinal Surgery Division, Armed Force Peng-Hu Hospital.

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Liver transplantation, Hepatobiliary surgery, Laparoscopic surgery

Managing immunosuppressants in liver transplantation – what have we learnt from COVID-19 pandemic?

Transplant activity has decreased during the COVID-19 pandemic. Adapted guidelines on potential deceased organ donors will be briefly mentioned.

Impaired immunity is a risk factor for critical outcome in transplant patients

with COVID-19. Factors that affect COVID-19 mortality in liver transplant patients and considerations for managing COVID-19 specific for organ transplantation are discussed herein.

How can we manage immunosuppression in such patients, choices of immunosuppressants and the possible underlying rationale will be elaborated. As for transplant patients receiving regular follow-up in outpatient clinic, innovative measures can be tried to minimize epidemiological exposure to COVID-19.

At last, guidelines from the American Association for the Study of Liver Diseases (AASLD) for managing post-transplant patients will be introduced.

移植年會議程

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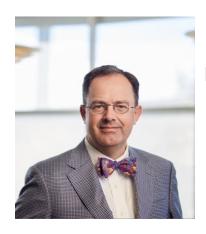
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Optimizing Immunosuppression: To prevent DSA/AMR and BKVN in Kidney Transplantation

Kidney transplantation induces an immune response to the graft that leads to rejection and graft loss. Over the last 60 years immunosuppression has evolved to the point where tacrolimus (Tac) and mycophenolic acid (MPA)-based therapy are considered the standard of care to effectively control the immune response and prevent rejection. However, while leading to prolonged graft survival the combination of Tac/MPA can result in off-target effects [i.e., GI toxicity, renal toxicity, and infections (e.g., BK virus nephropathy)] that leads to physician-guided drug minimization and/or patient nonadherence. This in turn results in increased rates of de novo donor specific antibody and biopsy proven acute rejection. The purpose of this lecture will be to review the data the supports Tac/MPA-based immunosuppression and discuss its optimal use to navigate the requirement to provide sufficient drug therapy to control the alloimmune response while avoiding overimmunosuppression, which leads to off-target effects.



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Sedative and Immunosuppressive Effects of Dexmedetomidine in Transplantation

Dexmedetomidine, an α 2-adrenergic receptor agonist, is used as an anti-anxiety medication. It also exerts a cholinergic effect, thereby reducing the release of TNF- α . We examined our patients who underwent living donor liver transplantation. A trend toward the improvement of hepatocyte injury along with better liver function was observed in the dexmedetomidine-treated group during the first postoperative week. Subsequently, we generated a series of mouse models to investigate the effect of dexmedetomidine on sedationbased tolerance post-transplantation. Indeed, dexmedetomidine inhibited the proliferation of T cells and TNF- α production in a dosedependent manner. We used dexmedetomidine to treat skintransplanted mice and observed a significantly prolonged graft survival in mice that were administered a higher dose of dexmedetomidine. These results revealed that dexmedetomidine exerts a dual effect of sedation and immunosuppression. This light-sedation approach will not only make patients calmer in the intensive care unit but also protect allografts from injury. The link between sedation and immunity may be designed toward therapeutic manipulation of the immune response.



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Long-Term Outcomes of Living Kidney Donors: Lessons Learned From Taiwanese Nationwide Cohort

The prevalence and incidence of hemodialysis are still quite high in Taiwan. Kidney transplantation could provided better life quality and survival benefit for patients with end stage kidney disease. However, shortage of kidney donor is always a major issue. Beyond cadaveric kidney transplantation, living kidney transplantation donated from recipient's relatives is the alternative choice. Safety issue is most important concern for living kidney donor. It is very important to fully understand the risk for living kidney donor and share the information with all stockholders, including nephrologists, transplant surgeons, transplant team members, potential donor, recipients and their families. In this talk, we will discuss the risk of end stage renal failure and death after kidney donation from literature review and the analysis of data of 1232 living kidney donor from Taiwan national health insurance research database.



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Peri-transplant treatment strategies for HCC beyond UCSF criteria

According to 2020 update on the management consensus guideline for hepatocellular carcinoma (HCC) by the Taiwan Liver Cancer Association and Gastroenterological Society of Taiwan, HCC is one of the leading causes of cancerrelated death in Taiwan. Curative treatment for HCC is limited and liver transplantation (LT), in theory, gives the best option for cure for patients with HCC with poor liver function not eligible for surgical resection or local ablation because it provides the widest tumor free margin while replacing diseased liver parenchyma with healthy liver tissue. Milan criteria has been served as the benchmark for patient selection in HCC undergoing LT, however, it restricted only to limited number of patients that fulfill the criteria for LT when they were diagnosed. Numerous modified less strict criterias, such as UCSF or up-to-seven criteria, have been speculated for patient selection that could achieve comparable outcomes to those consistently within the Milan criteria. For those HCCs beyond UCSF criteria, lots of peritransplant treatment strategies have been employed as a downstaging strategy to recruit more HCC patients for inclusion into the LT waiting list. These modalities include salvage liver resection, radiofrequency ablative therapy, trans-arterial chemo-embolization, radio-embolization, stereotactic body radiotherapy, or their combination. In the present talk, the last evidence of efficacy in those HCCs that beyond criteria who undergo different downstaging therapies will be reviewed.

For advanced stage HCCs, recent published cases on immune checkpoint inhibitor (ICI) for downstaging followed by LT have shown promising results. However, reports have also revealed immunotherapy to induce graft loss following LT. In this presentation, the first preliminary case reports illustrate ICI for downstaging therapy of advanced HCC before LT will also be introduced.





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