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臨床科別：過敏免疫風濕科

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專長

僵直性脊椎

自體免疫疾病

風濕關節炎

紅斑狼瘡

免疫

臨床試驗

痛風

整合醫學

過敏

學歷

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經歷

台中榮總家庭醫學科醫師
高雄榮總過敏免疫風濕科總醫師
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中山醫學大學附設醫院人體試驗委員會主任委員
行政院衛生署人體試驗/研究倫理委員會訪查委員

臨床試驗(執行中)

1. 為期 52 週的隨機、雙盲、安慰劑對照研究，目的在於評估活性自體抗體呈陽性的全身性紅斑狼瘡成年患者使用 Belimumab 後應特別注意的不良事件

2. 一項以患有類風濕性關節炎之患者為對象，評估 baricitinib 長期安全性暨療效之第三期多中心試驗
3. 使用TOFACITINIB (CP-690,550) 治療乾癬性關節炎的長期、開放、延伸性試驗
4. 評估兩種劑量的Tofacitinib與一種腫瘤壞死因子抑制劑 (TNFi) 用於類風濕性關節炎受試者的第 3B/4 期隨機分配安全性指標試驗
5. 以大數據研究免疫風濕疾病及藥物之長期效益以及風險
6. SpA登錄計畫及免疫基因體研究
7. 微生物菌相與唾液蛋白質組學在臨床診斷的應用與探討
8. 以人工智慧深度學習建立免疫風濕疾病影像自動判讀系統
9. 一項隨機分配、雙盲、安慰劑對照驗證試驗，在對MTX反應不足或無法耐受的類風濕性關節炎(RA)患者中，評估ASP015K的安全性與療效
10. 篩選帶有HLA同合子之細胞捐贈者
11. 風濕免疫疾病影響子代健康- 一項全國大數據的觀察型研究
12. 中軸型脊椎關節炎前瞻世代研究分類
13. 台灣中重度風濕性關節疾病患者，生物製劑(TNF inhibitor)中斷使用之流病分析，2010-2017年全國健保資料庫研究
14. 一項第3期、多中心、隨機分配、雙盲、安慰劑與活性藥物對照試驗，針對中度至重度斑塊型乾癬的受試者評估以Guselkumab治療的療效與安全性
15. 一項多中心、隨機分配、雙盲、安慰劑對照的24週試驗接續長期評估使用Ixekizumab (LY2439821) 於經歷生物疾病修飾抗風濕病藥物 (bDMARD) 的活性乾癬性關節炎 (PsA) 病患之療效與安全性
16. 針對充分達到24週反應的無X光異常之早期軸心型脊椎關節炎受試者，評估停止接受與再度接受ETANERCEPT治療的一項多中心、開放性試驗
17. 一項第 3 期、多中心、隨機分配、安慰劑對照的雙盲試驗，在罹患活動性軸心型脊椎關節炎 (AxSpA) 且沒有僵直性脊椎炎 (AS) 之 X 光證據及客觀發炎徵象的受試者中，評估 Certolizumab Pegol 的療效及安全性
18. 一項第三期、隨機分配、雙盲之臨床試驗，比較ABT-494、Adalimumab與安慰劑於接受穩定劑量之Methotrexate(MTX)治療未獲良好控制之中重度活動性類風濕性關節炎患者之安全性與療效

19. 一項隨機分配、雙盲、平行分組、多中心試驗，在抗腫瘤壞死因子 (TNF) 療法療效反應不足且正在接受背景methotrexate (MTX) 療法的中至重度類風濕性關節炎 (RA) 病患中，比較 JHL1101與在歐盟上市的MabThera®的藥物動力學、藥效學、免疫原性、安全性及療效
20. 一項隨機分配、雙盲、安慰劑和活性藥物對照、多中心、第三期試驗，在罹患中至重度活動性類風濕性關節炎，且對 Methotrexate 療效反應不足的受試者中，評估 Filgotinib 與 Methotrexate 併用 52 週的療效及安全性
21. 以軸心型脊椎關節炎病患評估Ixekizumab (LY2439821) 療效維持情況之一項104週多中心、長期延伸試驗 (包括一個雙盲、安慰劑對照40週隨機分配退出-再度治療期)
22. 一項第三期多中心、隨機分配、雙盲、安慰劑對照及開放性延伸試驗，評估KHK4827 對軸心型脊椎關節炎受試者之療效與安全性
23. 一項第三期、多中心、隨機分配、雙盲、安慰劑對照試驗，於患有活動性乾癬性關節炎的受試者，包括曾接受抗腫瘤壞死因子 (TNF α)生物製劑治療者，評估皮下給予Guselkumab之療效和安全性
24. 針對曾對至少一種非生物性疾病修飾型抗風濕藥物(DMARD)反應不佳的活動性乾癬性關節炎受試者中，對Upadacitinib(ABT-494) 與安慰劑以及Adalimumab進行比較的一項第三期、隨機分配、雙盲試驗 - SELECT –PsA 1
25. 一項多中心、雙盲、長期延伸試驗，在罹患類風濕性關節炎的受試者中，評估 Filgotinib 的安全性和療效
26. 一項針對在台灣接受Adalimumab療法的僵直性脊椎炎病患探討臨床反應的真實世界、前瞻性、觀察性試驗
27. 一項隨機分配、雙盲、安慰劑對照的多機構合作臨床試驗，於誘導劑量期之後進行隨機停藥維持劑量期，以評估Mirikizumab在中度至重度斑塊型乾癬病患中的療效與安全性
28. 隨機、部分盲性、活性藥物對照的多中心試驗，評估 secukinumab 用於活動性僵直性脊椎炎患者，治療 104 週相較於 GP2017 (adalimumab 生技仿製藥) 的放射影像惡化減少之療效，以及持續 2 年的長期安全性、耐受性及療效
29. 一項隨機分配、雙盲、安慰劑對照的第二期試驗，在患有全身性紅斑性狼瘡的受試者中，評估 BMS-986165 的療效及安全性

30. 一項多中心、隨機分配、雙盲、安慰劑對照、平行分組，使用 Ustekinumab 於活動性全身性紅斑性狼瘡受試者的試驗
31. 脊椎關節炎的週邊關節侵犯之評估研究
32. 一項多中心、長期延伸試驗，評估 Mirikizumab 於中度至重度斑塊型乾癬病患的長期安全性與維持治療效果
33. 針對曾對於至少一種疾病調節抗風濕藥物(DMARD)治療反應不佳或無法耐受的活動性乾癬性關節炎 (PsA) 受試者使用 Risankizumab 與安慰劑進行比較之第3期、隨機分配、雙盲性試驗
34. 一項多中心試驗，針對患有全身性紅斑性狼瘡的受試者，描述 BMS-986165 的長期安全性及療效特性
35. 一項第 2B 期、雙盲、隨機、安慰劑對照、多中心、劑量範圍試驗，旨在評估 PF-06700841 在活動性全身性紅斑性狼瘡 (SLE) 患者中的療效和安全性

臨床試驗(已結案)

1. 僵直性脊椎炎之中藥方劑療效評估
2. 開放性、多中心臨床試驗以評估台灣慢性乾癬症病患接受每週 肌肉注射 AMEVIVE 一個療程後之療效及安全性
3. 中藥方劑治療僵直性脊椎炎之先導性臨床試驗
4. 以基因微陣列法研究「僵直性脊椎炎」之基因表達
5. 喜樂美食品治療高尿酸血症病人之先導性研究
6. 以『自律神經分析儀』探討僵直性脊椎炎與其他免疫疾病病人自律神經功能之研究
7. Moducare® 治療過敏性鼻炎之隨機雙盲安慰劑對照式臨床試驗
8. 沙利竇邁治療結節性癢疹之療效評估計畫
9. 奈米珍珠鈣抗氧化延緩衰老及增進鈣生物利用率
10. AGTC-179 治療風濕病之隨機雙盲安慰劑控制式之臨床試驗
11. 非侵入性研究僵直性脊椎炎的基因多型性相關研究
12. 中藥方劑治療高尿酸血症之先導性試驗
13. 破骨細胞與內質網壓力在僵直性脊椎炎致病機制中所扮演的角色
14. MORA 生物能共振檢測儀用於過敏病患之臨床試驗
15. 以 etanercept 治療僵直性脊椎炎的多中心、開放性研究
16. 以高頻熱凝療法阻斷腰椎脊神經內側枝治療腰椎小面關節症候群
17. 國際性、多中心研究利用僵直性脊椎炎療效評估標準量表作為脊椎關節炎病患分類標準

18. 抗腫瘤壞死因子(enbrel)對僵直性脊椎炎血清中TLR, BMP, TARIL, MMP及多種cytokines(TNF, IL18, IL6, IL8, IL10, IL1)等之影響
19. Pioglitazone hydrochloride 30mg 錠劑生體相等性試驗
20. 比較Etoricoxib 與 Aceclofenac 對類風濕性關節炎患者的療效
21. 中藥方劑治療高尿酸血症之先導性試驗
22. 一個開放性的試驗以評估MabThera® (rituximab)用於治療對一種或多種抗腫瘤壞死因子藥物反應不佳或耐受性不佳的類風濕性關節炎發作病患
23. 痛風散治療高尿酸血症之先導性試驗
24. 免疫調節草本複方食品莊松榮精氣神茶飲之隨機雙盲安慰劑控制式臨床試驗
25. 一個自第II至III階段、多中心、隨機、雙盲、以安慰劑對照之臨床研究。Abatacept相對於安慰劑對正在接受Mycophenolate Mofetil 及Glucocorticosteroids治療因全身紅斑性狼瘡引起活動性增生性腎絲球體腎炎之受試者其療效和安全性的評估
26. 一項第三期、多中心、隨機分配、雙盲、安慰劑對照、52週的研究，評估全人類 BlyS 單株抗體 Belimumab (HGS1006, LymphoStat-B™) 對於全身性紅斑狼瘡(SLE) 受試者的療效和安全性
27. Health Baby裝置之心率變異分析與體溫功能評估研究
28. Tramadol/Acetaminophen (Ultracet®)藥劑附加治療於僵直性脊椎炎病患
29. 比較Etanercept 與Adalimumab治療於僵直性脊椎炎病患
30. 第三階段、多中心、隨機分派、雙盲、雙啞 之對照研究，比較 Abatacept以皮下注射與靜脈注射用於正在使用Methotrexate治療類風濕性關節炎但對Methotrexate反應不足的患者，其療效與安全性之評估
31. 丹參附加西藥於台灣原發性高血壓病患之有效性與耐受性—十二週之隨機、雙盲、安慰劑控制式臨床試驗
32. 評估Thalidomide治療合併有關節炎或無關節炎之斑塊尋常性乾癬病患之有效性與安全性 - 二十四週之隨機、雙盲、安慰劑控制式臨床試驗
33. 一項多中心，將全人類 BlyS 單株抗體 Belimumab (HGS1006,LymphoStat-B™)用於完成第三期 HGS1006-C1056 或 HGS1006-C1057 試驗計畫之全身性紅斑狼瘡 (SLE) 受試者的延續試驗
34. 評估樟芝輔助改善類風濕性關節炎之有效性與安全性 - 一項為期十二週、隨機、雙盲、安慰劑、控制式臨床試驗

35. 計畫性細胞死亡-1及其配位基與第四類細胞毒殺性T淋巴抗原基因多形性與台灣僵直性脊椎炎之相關性
36. 評估保健食品Domilex輔助改善類風濕性關節炎之隨機雙盲安慰劑控制式臨床試驗
37. 併用疾病修飾性抗風濕病藥物(DMARD)治療活動期類風濕性關節炎的病人，接受LY2439821(抗IL-17抗體)多劑皮下注射的第2期劑量變動試驗
38. 一項隨機、開放、對照式臨床試驗評估抗腫瘤壞死因子生物製劑治療僵直性脊椎炎患者合併使用非類固醇消炎止痛藥之安全性與有效性
39. 紅麴加上益生菌對降血脂作用之評估-生化分析及機轉研究
40. 中藥複方OA 1改善骨關節炎症狀之人體試驗 - 一項為期十二週之隨機雙盲安慰劑對照式臨床試驗
41. 建立以醫院為基礎的僵直性脊椎炎病患世代，並且探討骨骼重塑與自體免疫耐受性的可能效應
42. 雪樟芝輔助改善C型肝炎之十二週隨機雙盲安慰劑控制式臨床試驗
43. 一項多中心、12週、雙盲、安慰劑對照、隨機分配的試驗，以恩博併用非類固醇消炎止痛藥治療無影像學變化之中軸脊椎關節炎成年患者，並進行為期92週開放性延伸研究
44. TNF- α 、TNF- α 接受器與microRNA-146a基因多形性與僵直性脊椎炎發生之相關
45. 一項第III期、2階段、隨機分配、雙盲、安慰劑對照、多中心臨床試驗，評估2種MK-0663/Etoricoxib劑量用於類風濕性關節炎病患時的相對療效和耐受性
46. 一項第III期、2階段、隨機分配、雙盲、安慰劑對照、多中心臨床試驗，評估2種MK-0663/Etoricoxib劑量用於類風濕性關節炎病患時的相對療效和耐受性
47. 在僵直性脊椎炎病患中測試MK-0663/Etoricoxib2種劑量之相對療效和耐受性的一項第III期、2階段、隨機分配、雙盲、活性對照、多中心臨床試驗
48. 一個採隨機、安慰劑控制評估Tocilizumab合併使用Methotrexate在治療中度至重度的類風濕性關節炎患者之臨床試驗
49. 一項針對活動性乾癬性關節炎受試者評估使用兩種Apremilast (CC-10004)劑量的第3期、多中心、隨機分配、雙盲、安慰劑對照、平行分組之療效與安全性試驗

50. 一項第三期、多中心、隨機分配、雙盲、安慰劑對照試驗，評估全身性紅斑性狼瘡(SLE)病患使用皮下注射LY2127399的療效與安全性
51. 一項評估全身性紅斑性狼瘡受試者使用A-623的療效、安全性和耐受性之隨機雙盲第2b期試驗
52. 表面擴散係數分析在修格連氏症候群腮腺之研究
53. 隨機分組、雙盲、以安慰劑對照的劑量範圍決定試驗：針對先前以TNF 阻斷劑治療無效的活躍性類風濕性關節炎病患，評估以皮下注射方式使用CDP6038治療12週的療效與安全性。第二期
54. 隨機、雙盲、安慰劑對照的多機構合作試驗，研究以secukinumab 治療 16 週的療效，並持續 2 年評估活動性僵直性脊椎炎患者的長期安全性、耐受性與療效
55. 一項第三期、多中心、隨機分配、雙盲、安慰劑對照的平行試驗，評估以兩種口服CP-690,550劑量治療中度至重度慢性斑塊型牛皮癬患者的療效及安全性
56. 紅景天輔助改善慢性阻塞性肺病之隨機雙盲安慰劑控制式臨床試驗及機轉
57. 一項第三期、多中心、開放性試驗，評估以兩種口服CP-690,550劑量治療中度至重度慢性斑塊型牛皮癬患者的長期安全性及耐受性
58. 這是一項第2期、多中心、開放標示、追蹤試驗，用以評估完成RA0083試驗活動性類風濕性關節炎的亞洲受試者，以皮下注射接受CDP6038的長期安全性與療效
59. 一項開放性、長期的安全性延伸試驗，針對已完成AN-SLE3321 (PEARL-SC)試驗計劃之全身性紅斑性狼瘡受試者
60. (OSKIRA-Asia-1X)：在亞洲進行的一項長期試驗，評估Fostamatinib用於治療類風濕性關節炎的安全性
61. 一項針對全身性紅斑性狼瘡 (Systemic Lupus Erythematosus) 受試者評估MEDI-546之療效與安全性的第二期、隨機分組試驗
62. 一項第3b期、多中心、開放試驗，評估紅斑性狼瘡 (SLE) 病患接受LY2127399皮下注射劑之長期療效與安全性 (ILLUMINATE - X)
63. 一項前瞻性、隨機、雙盲、安慰劑對照、平行性、多中心、第三期試驗，評估ENIA11併用Methotrexate與單用Methotrexate治療類風濕性關節炎病患的療效性與安全性
64. 一項前瞻性、隨機、雙盲、安慰劑對照、平行性、多中心、第三期試驗，評估ENIA11併用Methotrexate與單用Methotrexate治療類風濕性關節炎病患的療效性與安全性

65. 一項對接受DMARD藥物治療後仍有活動性類風濕性關節炎的受試者以皮下注射給予CNTO 136 (sirukumab, 一種人類抗介白素6 [IL-6]單株抗體)的多中心、隨機分配、雙盲、安慰劑對照、平行分組之試驗
66. 一項對接受抗腫瘤壞死因子 α (Anti-TNF α)療法後仍有活動性類風濕性關節炎的受試者以皮下注射給予CNTO 136 (sirukumab, 一種人類抗介白素6 [IL-6]單株抗體)的多中心、隨機分配、雙盲、安慰劑對照、平行分組之試驗
67. 一項以患有中度到重度活動性類風濕性關節炎, 且經methotrexate 治療後反應不佳之患者為對象, 評估接受baricitinib 療法所得療效與安全性之隨機、雙盲、安慰劑暨活性對照第三期試驗
68. 此多國、橫斷式研究乃為了解全身慢性免疫疾病病患, 對於服藥用治療順從性的一般與特殊認知
69. 評估AC-201用於接受降尿酸藥治療的痛風患者之隨機、雙盲、安慰劑對照之二期臨床試驗
70. 隨機分配、雙盲、安慰劑對照之第三期臨床試驗, 評估Baricitinib (LY3009104) 用於使用傳統疾病修飾抗類風濕藥物反應不佳之中度至重度活動性類風濕性關節炎患者之療效與安全性
71. 一項評估非生物性 DMARD 療法中加入 sarilumab (用於接受TNF- α 拮抗劑但無獲得充分緩解或無法耐受之類風濕性關節炎患者) 的療效與安全性之隨機分配、雙盲、平行組、安慰劑對照研究
72. 藥物基因體學分析強直性脊柱炎患者TNF- α 受體阻滯劑治療的反應偵測
73. 脊椎關節炎之合併症流行病學研究
74. 102年度「教學醫院成立中藥臨床試驗中心」
75. 一項評估Blisibimod用於全身性紅斑狼瘡受試者之療效與安全性的隨機分配、雙盲、安慰劑對照、第3期試驗
76. 評估 Tofacitinib 治療患有活動性僵直性脊椎炎 (AS) 受試者的療效及安全性, 在一項第二期、隨機分配、雙盲、安慰劑對照、劑量範圍的研究
77. 骨質疏鬆症的藥物使用模式、治療滿意度和控制不力研究 (MUSIC-OS): 亞太地區停經後婦女在骨質疏鬆症治療方面的臨床實務研究
78. 微型核糖核酸-125b及其標的基因TNF- α 與僵直性脊椎炎之相關

79. 一項第 3 期、隨機分組、雙盲、安慰劑控制的試驗，評估 2 種劑量的 Tofacitinib (CP-690,550) 或 Adalimumab 對於活動性乾癱性關節炎病患的療效及安全性
80. 生物製劑減量及暫緩續用對於類風濕性關節炎患者的臨床表現及醫療資源使用情況影響的回溯性病歷研究
81. 評估 secukinumab 治療活動性僵直性脊椎炎患者之臨床效益持續性、安全性、及耐受性之延伸試驗
82. 評估新興市場的風濕門診中，早期無X光中軸型脊椎關節炎(non-radiographic Axial SpA)在慢性發炎性背痛病患的盛行率之橫斷式前瞻性非介入性之流行病學研究
83. 一項針對已於 CNTO136ARA3002 (SIRROUND-D) 與 CNTO136ARA3003 (SIRROUND-T) 試驗中完成治療的類風濕性關節炎受試者研究CNTO136 (sirukumab)長期安全性與療效的多中心、平行分組試驗
84. 在僵直性脊椎炎中，循環微型核糖核酸 (miR-27, miR-29, miR-125b, miR-146a, miR-155) 與其標的基因、發炎體基因多形性、骨重塑與發炎標記、以及臨床表徵之相關
85. 一項以患有僵直性脊椎炎的受試者為對象，採用三種劑量、以皮下方式給予藥物BI 655066，用以證明概念及探索劑量，且為期 48 週的第二期、隨機分配、雙盲、安慰劑對照試驗
86. 一項第 3 期、隨機分配、雙盲、安慰劑對照的試驗，評估 2 種劑量的 TOFACITINIB (CP-690,550)在活動性乾癱性關節炎及對至少一種 TNF 抑制劑反應不足之受試者的療效與安全性
87. 一項第 3b /4 期隨機分配、雙盲的試驗，在中度至嚴重活動性類風濕性關節炎受試者中，比較5 毫克劑量的 tofacitinib 併用及不併用 methotrexate，與 adalimumab 併用 methotrexate 的研究
88. 一項第三期隨機分配、雙盲試驗，針對曾對於METHOTREXATE 反應不佳的中度至重度活動性類風濕性關節炎受試者，評估PF-06410293與ADALIMUMAB併用METHOTREXATE的療效和安全性
89. 以健保資料庫分析台灣常見風濕病之全國流行病學調查及共病研究
90. 教學醫院推動中藥臨床試驗療效評估
91. 口腔微生物菌相及唾液蛋白質組學在口腔乾燥診斷上的應用
92. 低劑量電腦斷層在僵直性脊椎炎之應用

93. 一項第3期、多中心、隨機分配、雙盲試驗，針對中度至重度斑塊型乾癬且對 Ustekinumab 反應不佳的受試者評估以 Guselkumab 治療的療效與安全性
94. 一項使用 Ustekinumab 治療活動性全身性紅斑性狼瘡病患的多中心、隨機分配、雙盲、安慰劑對照、概念驗證試驗
95. 一項針對未曾使用抗腫瘤壞死因子(TNF α)藥物之活動性放射影像軸心型脊椎關節炎受試者，評估 Ustekinumab 療效和安全性的第三期、多中心、隨機分配、雙盲、安慰劑對照試驗
96. 一項針對具有抗腫瘤壞死因子(TNF α)藥物頑抗性之活動性放射影像軸心型脊椎關節炎受試者，評估 Ustekinumab 療效和安全性的第三期、多中心、隨機分配、雙盲、安慰劑對照試驗
97. 一項針對活動性無放射影像異常之早期軸心型脊椎關節炎受試者評估 Ustekinumab 療效和安全性的第三期、多中心、隨機分配、雙盲、安慰劑對照試驗
98. ASAS 健康量表之國際信效度研究計畫
99. 一項隨機分配、雙盲、安慰劑對照確認試驗，評估 ASP015K 用於疾病修飾型抗風濕藥物 (DMARD) 治療反應不佳的類風濕性關節炎 (RA) 患者之安全性與療效
100. 多中心、開放性(A部分)後進行隨機分配、雙盲、平行分組、安慰劑對照試驗(B部分)，針對患有活性軸心型脊椎關節炎(axSpA)的受試者，接受 Certolizumab pegol 200 mg 每2週一次(Q2W) 或 200 mg 每4週一次(Q4W)，相較於安慰劑治療，評估維持緩解的效果
101. 一個第IIb期、隨機、雙盲、安慰劑對照研究，評估干擾素基因標記的中和作用，以及 IFN α -Kinoid 對全身性紅斑狼瘡成人患者的臨床療效標記
102. Ixekizumab (LY2439821) 使用於未曾接受 bDMARD 之放射線影像異常軸心型脊椎關節炎病患的一項多中心、隨機分配、雙盲、活性藥物與安慰劑對照 16 週試驗與安全性及療效之長期評估追蹤
103. 一項隨機、雙盲、雙虛擬、多中心、活性對照試驗，在患有潰瘍性結腸炎的受試者中，評估靜脈注射型 Vedolizumab 相較於皮下注射型 Adalimumab 的療效及安全性
104. 針對 Baricitinib 在全身紅斑性狼瘡 (Systemic Lupus Erythematosus, SLE) 病患的隨機分配、雙盲、安慰劑對照、平行分組、第2期試驗
105. 一項針對健康受試者的第1期、隨機分配、雙盲、安慰劑對照、單劑遞增劑量試驗，以及針對輕度至中度全身性紅斑性狼瘡受試者的 JNJ-55920839 多重劑量試驗 - B 部分

- 106.第2期、隨機分配、多中心、雙盲、劑量範圍、安慰劑對照的調整設計試驗，針對罹患中度至重度類風濕性關節炎且對Methotrexate併用或不併用腫瘤壞死因子(TNF)抑制劑反應不足的受試者，評估BMS-986142的療效與安全性/藥物動力學
- 107.一項在活動性乾癱性關節炎患者中探討BI 655066/ABBV-066/risankizumab的隨機、雙盲、安慰劑對照、概念驗證、劑量範圍試驗
- 108.探討藉由ASAS提出11項心理測量評估中軸型脊椎關節炎病發之特徵研究
- 109.一項第3期、隨機分配、雙盲、活性藥物對照、多中心試驗，於腕關節或膝關節骨關節炎的受試者皮下注射tanezumab長期安全性與療效研究
- 110.ASP015K 延伸試驗–開放標記延伸試驗，對象為完成 ASP015K 第 IIb 期或第 III 期試驗的類風濕性關節炎患者
- 111.針對接受全膝關節、全腕關節或全肩關節置換術的Tanezumab試驗受試者所進行的一項第3期、多中心、長期觀察性試驗
- 112.一項隨機分配、雙盲、安慰劑和活性藥物對照、多中心、第三期試驗，在罹患中至重度活動性類風濕性關節炎，且未曾接受Methotrexate (MTX) 療法的受試者中，評估單獨使用 Filgotinib 與併用 MTX 52 週的療效及安全性
- 113.針對已於試驗1311.5中完成第24週回診之乾癱性關節炎受試者，研究Risankizumab安全性的一項第2期、單組、開放性延伸試驗
- 114.以單一劑量TLC599注射退化性膝關節炎病患之一項第IIa期、隨機分配、雙盲、安慰劑對照、劑量探索臨床試驗

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