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# CLINICAL TRIALS FOR FSGS PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE | DRUG/<br>COMPOUND          | EGFR<br>IN ML/MIN                         | PHASE |
|--|--|--------------|----------------------------|---|-------|
| <b>EPPIK</b><br>(Travere)  | FSGS, MCD, IgAN,<br>IgAV, Alport Syndrome                                    | 1-17         | Sparsentan                 | ≥ 30                                      | 2     |
| <b>DUPLEX (ENROLLMENT COMPLETE)</b><br>(Travere)   | FSGS   | 8-75         | Sparsentan                 | ≥ 30                                      | 3     |
| <b>THE PODO TRIAL</b><br>(Pfizer)  | FSGS   | 18+          | PF-06730512                | ≥ 30                                      | 2     |
| <b>THE TRACTION-2 STUDY</b><br>(Goldfinch Bio)   | FSGS, treatment-<br>resistant MCD, Lipoid<br>Nephrosis, Diabetes<br>Mellitus | 18-75        | GFB-887                    | ≥ 30                                      | 2     |
| <b>STUDY OF VX-147 IN APOL1-MEDIATED PROTEINURIC<br/>KIDNEY DISEASE</b><br>(Vertex)  | Proteinuric Kidney<br>Disease  | 18-60        | VX-147                     | N/A                                       | 2/3   |
| <b>STUDY OF VX-147 IN APOL1-MEDIATED KIDNEY DISEASE</b><br>(Vertex)  | APOL1-Mediated FSGS  | 18-65        | VX-147                     | ≥ 30                                      | 2     |
| <b>AFFINITY</b><br>(Chinook)   | FSGS, IgAN, Alport<br>Syndrome, DKD,<br>Diabetic Nephropathy<br>Type 2       | 18+          | Atrasentan                 | ≥ 30                                      | 2     |
| <b>STUDY OF BI 764198 IN FSGS</b><br>(Boehringer Ingelheim)  | FSGS   | 18-75        | BI 764198                  | ≥ 30                                      | 2     |
| <b>POST-APPROVAL STUDY OF LIPOSORBER LA-15 SYSTEM FOR<br/>PEDIATRIC &amp; DRUG RESISTANT ADULT PATIENTS WITH FSGS</b><br>(Kaneka)    | Post-transplant recurrent<br>FSGS or Primary FSGS                            | 5-75         | Liposorber LA-15<br>System | ≥ 45 or post-<br>transplant<br>recurrence | N/A   |
| <b>TUMOR NECROSIS FACTOR INHIBITION IN FSGS AND<br/>TREATMENT-RESISTANT MCD</b><br>(University of Michigan)                          | FSGS, MCD  | 6-70         | adalimumab                 | > 45                                      | 2     |
| <b>STUDY OF VB119 IN STEROID-SENSITIVE PRIMARY MCD OR<br/>PRIMARY FSGS</b><br>(ValenzaBio)   | FSGS, MCD  | 18+          | VB119                      | ≥ 60                                      | 2     |
| <b>STUDY OF ANG-3070 IN PRIMARY GLOMERULAR DISEASE</b><br>(Angion)   | Glomerular Disease   | 18+          | ANG-3070                   | ≥ 40                                      | 2     |
| <b>STUDY OF BI 690517 ALONE AND IN COMBINATION WITH<br/>EMPAGLIFLOZIN IN DIABETIC AND NON-DIABETIC CKD</b><br>(Boehringer Ingelheim) | Chronic Kidney Disease   | 18+          | BI 690517                  | ≥ 30 &<br>< 90                            | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI 685509<br/>ON KIDNEY FUNCTION IN CKD WITHOUT DIABETES</b><br>(Boehringer Ingelheim) | Chronic Kidney Disease   | 18+          | BI 685509                  | ≥ 20 &<br>< 90                            | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)  | Chronic Kidney Disease   | 18+          | AZD5718,<br>Dapagliflozin  | 20-75                                     | 2     |

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# CLINICAL TRIALS FOR FSGS PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE            | DRUG/<br>COMPOUND                       | EGFR<br>IN ML/MIN | PHASE |
|--|--|-------------------------|---|-------------------|-------|
| <b>ZENITH-CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease   | 18+                     | Zibotentan<br>Dapagliflozin             | ≥ 20              | 2     |
| <b>EAGLE</b><br>(Reata)  | Chronic Kidney Disease,<br>Alport Syndrome                           | 12+                     | Bardoxolone<br>methyl                   | ≥ 20              | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI<br/>OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE<br/>WITH CKD</b><br>(Bayer)   | Chronic Kidney Disease   | 6 months<br>to 17 years | Finerenone<br>(Kerendia,<br>BAY94-8862) | ≥ 30              | 3     |
| <b>FIND-CKD</b><br>(Bayer)   | Chronic Kidney Disease   | 18+                     | Finerenone<br>(BAY94-8862)              | ≥ 25 and < 90     | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD<br/>STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON<br/>DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi)                              | Secondary<br>Hyperparathyroidism<br>Chronic Kidney Disease           | 5-18                    | Doxercalciferol<br>(GZ427397)           | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)   | Chronic Kidney Disease<br>Stage IV                                   | 18-80                   | US-APR2020                              | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS AND<br/>SAFETY OF MK-2060 IN PATIENTS WITH ESRD RECEIVING<br/>HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)                               | End Stage Renal Disease<br>& Kidney Failure                          | 18+                     | MK-2060                                 | N/A               | 2     |
| <b>FREEDOM-1</b><br>(Talaris)  | De novo living donor<br>transplant                                   | 18+                     | FCR001                                  | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)  | De novo living donor<br>transplant (3-12 months<br>after transplant) | 18+                     | FCR001                                  | N/A               | 2     |
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY<br/>REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT<br/>RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant   | 18-75                   | Extracorporeal<br>photopheresis         | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS<br/>IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor<br>transplant                                   | 18-65                   | Apheresis                               | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF<br/>ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT<br/>(ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant   | 18-70                   | VIB4920 &<br>Belatacept                 | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH<br/>BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28<br/>&amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor<br>transplant                                   | 18-70                   | CD28 & IL-6<br>Receptor<br>Antagonists  | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant<br>rejection                                       | 18-70                   | TX200-TR101                             | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR MCD PATIENTS

| TRIAL NAME<br>(sponsor)   | PATIENT<br>DIAGNOSIS  | AGE<br>RANGE            | DRUG/<br>COMPOUND                       | EGFR<br>IN ML/MIN | PHASE |
|---|---|-------------------------|---|-------------------|-------|
| <b>EPPIK</b><br>(Traverse)  | MCD, FSGS, IgAN,<br>IgAV, Alport Syndrome                                   | 1-17                    | Sparsentan                              | ≥ 30              | 2     |
| <b>THE TRACTION-2 STUDY</b><br>(Goldfinch Bio)  | Treatment-resistant<br>MCD, FSGS, Lipoid<br>Nephrosis, Diabetes<br>Mellitus | 18-75                   | GFB-887                                 | ≥ 30              | 2     |
| <b>TUMOR NECROSIS FACTOR INHIBITION IN FSGS AND<br/>TREATMENT-RESISTANT MCD</b><br>(University of Michigan)   | Treatment-resistant<br>MCD, FSGS  | 6-70                    | adalimumab                              | > 45              | 2     |
| <b>STUDY OF VB119 IN STEROID-SENSITIVE PRIMARY MCD OR<br/>PRIMARY FSGS</b><br>(ValenzaBio)  | MCD, FSGS   | 18+                     | VB119                                   | ≥ 60              | 2     |
| <b>STUDY OF ANG-3070 IN PRIMARY GLOMERULAR DISEASE</b><br>(Angion)  | Glomerular Disease  | 18+                     | ANG-3070                                | ≥ 40              | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI 685509<br/>ON KIDNEY FUNCTION IN CKD WITHOUT DIABETES</b><br>(Boehringer Ingelheim)                      | Chronic Kidney Disease  | 18+                     | BI 685509                               | ≥ 20 &<br>< 90    | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease  | 18+                     | AZD5718,<br>Dapagliflozin               | 20-75             | 2     |
| <b>EAGLE</b><br>(Reata)   | Chronic Kidney Disease,<br>Alport Syndrome                                  | 12+                     | Bardoxolone<br>methyl                   | ≥ 20              | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI<br/>OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE<br/>WITH CKD</b><br>(Bayer)                | Chronic Kidney Disease  | 6 months<br>to 17 years | Finerenone<br>(Kerendia,<br>BAY94-8862) | ≥ 30              | 3     |
| <b>FIND-CKD</b><br>(Bayer)  | Chronic Kidney Disease  | 18+                     | Finerenone<br>(BAY94-8862)              | ≥ 25 and<br>< 90  | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD<br/>STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON<br/>DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi) | Secondary<br>Hyperparathyroidism<br>Chronic Kidney Disease                  | 5-18                    | Doxercalciferol<br>(GZ427397)           | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)  | Chronic Kidney Disease<br>Stage IV  | 18-80                   | US-APR2020                              | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS AND<br/>SAFETY OF MK-2060 IN PATIENTS WITH ESRD RECEIVING<br/>HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)  | End Stage Renal Disease<br>& Kidney Failure                                 | 18+                     | MK-2060                                 | N/A               | 2     |
| <b>FREEDOM-1</b><br>(Talaris)   | De novo living donor<br>transplant  | 18+                     | FCR001                                  | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)   | De novo living donor<br>transplant (3-12 months<br>after transplant)        | 18+                     | FCR001                                  | N/A               | 2     |

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# CLINICAL TRIALS FOR MCD PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS            | AGE<br>RANGE | DRUG/<br>COMPOUND                | EGFR<br>IN ML/MIN | PHASE |
|--|---------------------------------|--------------|----------------------------------|-------------------|-------|
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant              | 18-75        | Extracorporeal photopheresis     | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor transplant | 18-65        | Apheresis                        | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant              | 18-70        | VIB4920 & Belatacept             | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor transplant | 18-70        | CD28 & IL-6 Receptor Antagonists | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant rejection     | 18-70        | TX200-TR101                      | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR IGAN PATIENTS

| TRIAL NAME<br>(sponsor)   | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE | DRUG/<br>COMPOUND | EGFR<br>IN ML/MIN | PHASE |
|---|--|--------------|-------------------|-------------------|-------|
| <b>EPPIK</b><br>(Travere)   | IgAN, FSGS, MCD,<br>IgAV, Alport Syndrome                              | 1-17         | Sparsentan        | ≥ 30              | 2     |
| <b>SPARTAN (LEICESTER, UK)</b><br>(Travere)   | IgAN   | 18+          | Sparsentan        | ≥ 30              | 2     |
| <b>PROTECT (ENROLLMENT COMPLETE)</b><br>(Travere)   | IgAN   | 18+          | Sparsentan        | ≥ 30              | 3     |
| <b>AFFINITY</b><br>(Chinook)  | IgAN, FSGS, Alport<br>Syndrome, DKD,<br>Diabetic Nephropathy<br>Type 2 | 18+          | Atrasentan        | ≥ 30              | 2     |
| <b>ALIGN</b><br>(Chinook)   | IgAN   | 18+          | Atrasentan        | ≥ 30              | 3     |
| <b>SAFETY &amp; TOLERABILITY OF BION-1301 IN HEALTHY<br/>VOLUNTEERS &amp; ADULTS WITH IGAN (CALIFORNIA, EL PASO<br/>TEXAS, LONDON)</b><br>(Chinook) | IgAN   | 18+          | BION-1301         | ≥ 30              | 1 & 2 |
| <b>VISIONARY</b><br>(Otsuka)  | IgAN   | 18+          | Sibeprenlimab     | ≥ 30              | 3     |
| <b>SAFETY &amp; EFFICACY OF VIS649 FOR IGAN (ENROLLMENT<br/>COMPLETE)</b><br>(Visterra)   | IgAN   | 18+          | VIS649            | ≥ 45              | 2     |
| <b>STUDY OF BI 690517 ALONE AND IN COMBINATION WITH<br/>EMPAGLIFLOZIN IN DIABETIC AND NON-DIABETIC CKD</b><br>(Boehringer Ingelheim)                | IgAN, FSGS, MN, DKD,<br>Hypertensive Kindey<br>Disease                 | 18+          | BI 690517         | ≥ 30 and<br>< 90  | 2     |
| <b>APPLAUSE-IGAN</b><br>(Novartis)  | IgAN   | 18+          | LNP023            | ≥ 20              | 3     |
| <b>ORIGIN</b><br>(Vera)   | IgAN, Berger Disease   | 18+          | Atacicept         | ≥ 30              | 2     |
| <b>NEFIGARD (ENROLLMENT COMPLETE)</b><br>(Calliditas)   | IgAN   | 18+          | Nefecon           | ≥ 35              | 3     |
| <b>THE DISCOVERY TRIAL (ENROLLMENT COMPLETE)</b><br>(Apellis)   | IgAN, MN, C3G,<br>Dense Deposit Disease,<br>or Lupus Nephritis         | 18+          | APL-2             | ≥ 30              | 2     |
| <b>SANCTUARY</b><br>(Alexion)   | IgAN, Lupus Nephritis  | 18-75        | Ravulizumab       | ≥ 30              | 2     |
| <b>STUDY OF ALXN2050 IN IGAN &amp; PROLIFERATIVE LUPUS<br/>NEPHRITIS</b><br>(Alexion)   | IgAN, Lupus Nephritis  | 18-75        | ALXN2050          | > 30              | 2     |

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# CLINICAL TRIALS FOR IGAN PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS                       | AGE<br>RANGE            | DRUG/<br>COMPOUND                       | EGFR<br>IN ML/MIN | PHASE |
|--|--|-------------------------|---|-------------------|-------|
| <b>TELITACEPT FOR INJECTION (RC18) IN IGAN</b><br>(RemeGen)  | IgAN                                       | 18-70                   | Telitacept                              | > 30              | 2     |
| <b>THE ARTEMIS-IGAN STUDY</b><br>(Omeros)  | IgAN                                       | 18+                     | Narsoplimab/<br>OMS721                  | ≥ 30              | 3     |
| <b>IGNAZ</b><br>(MorphoSys)  | IgAN                                       | 18-80                   | Felzartamab                             | UPCR:<br>≥ 1.0    | 2     |
| <b>STUDY OF CEMDISIRAN IN ADULTS WITH IGAN (ENROLLMENT COMPLETE)</b><br>(Alnylam)  | IgAN                                       | 18–65                   | Cemdisiran                              | ≥ 30              | 2     |
| <b>EFFECTIVENESS &amp; SAFETY OF IONIS-FB-LRX IN IGAN (AUSTRALIA, CANADA, NEW ZEALAND)</b><br>(Ionis)                              | IgAN                                       | 18–75                   | IONIS-FB-LRx                            | > 40              | 2     |
| <b>RENEW (ENROLLMENT COMPLETE)</b><br>(BioCryst)   | IgAN, C3G, MN                              | 18+                     | BCX9930                                 | ≥ 50              | 2     |
| <b>SAFETY AND EFFICACY OF AT-1501 IN IGA NEPHROPATHY</b><br>(Eledon)   | IgAN                                       | 18-99                   | AT-1501                                 | ≥ 30              | 2     |
| <b>SAFETY AND TOLERABILITY OF MEZAGITAMAB (TAK-079) IN IGAN WITH STABLE BACKGROUND THERAPY</b><br>(Takeda)                         | IgA Nephropathy,<br>IgAV                   | 18+                     | Mezagitamab                             | ≥ 45              | 1     |
| <b>STUDY OF ANG-3070 IN PRIMARY GLOMERULAR DISEASE</b><br>(Angion)   | Glomerular Disease                         | 18+                     | ANG-3070                                | ≥ 40              | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI 685509 ON KIDNEY FUNCTION IN CKD WITHOUT DIABETES</b><br>(Boehringer Ingelheim)   | Chronic Kidney Disease                     | 18+                     | BI 685509                               | ≥ 20 &<br>< 90    | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)  | Chronic Kidney Disease                     | 18+                     | AZD5718,<br>Dapagliflozin               | 20-75             | 2     |
| <b>ZENITH-CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease                     | 18+                     | Zibotentan<br>Dapagliflozin             | ≥ 20              | 2     |
| <b>EAGLE</b><br>(Reata)  | Chronic Kidney Disease,<br>Alport Syndrome | 12+                     | Bardoxolone<br>methyl                   | ≥ 20              | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE WITH CKD</b><br>(Bayer) | Chronic Kidney Disease                     | 6 months<br>to 17 years | Finerenone<br>(Kerendia,<br>BAY94-8862) | ≥ 30              | 3     |
| <b>FIND-CKD</b><br>(Bayer)   | Chronic Kidney Disease                     | 18+                     | Finerenone<br>(BAY94-8862)              | ≥ 25 and<br>< 90  | 3     |

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|--|--|--------------|--|-------------------|-------|
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi)                              | Secondary Hyperparathyroidism<br>Chronic Kidney Disease              | 5-18         | Doxercalciferol<br>(GZ427397)          | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)   | Chronic Kidney Disease<br>Stage IV                                   | 18-80        | US-APR2020                             | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS AND SAFETY OF MK-2060 IN PATIENTS WITH ESRD RECEIVING HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)                               | End Stage Renal Disease<br>& Kidney Failure                          | 18+          | MK-2060                                | N/A               | 2     |
| <b>FREEDOM-1</b><br>(Talaris)  | De novo living donor<br>transplant                                   | 18+          | FCR001                                 | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)  | De novo living donor<br>transplant (3-12 months<br>after transplant) | 18+          | FCR001                                 | N/A               | 2     |
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant   | 18-75        | Extracorporeal<br>photopheresis        | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor<br>transplant                                   | 18-65        | Apheresis                              | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant   | 18-70        | VIB4920 &<br>Belatacept                | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor<br>transplant                                   | 18-70        | CD28 & IL-6<br>Receptor<br>Antagonists | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant<br>rejection                                       | 18-70        | TX200-TR101                            | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR MEMBRANOUS PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE | DRUG/<br>COMPOUND           | EGFR<br>IN ML/MIN | PHASE |
|--|--|--------------|-----------------------------|-------------------|-------|
| <b>EFFICACY &amp; SAFETY OF LNP023 COMPARED WITH RITUXIMAB IN ADULTS WITH IDIOPATHIC MN</b><br>(Novartis)                        | MN   | 18+          | LNP023,<br>Rituximab        | ≥ 30              | 2     |
| <b>THE DISCOVERY TRIAL (ENROLLMENT COMPLETE)</b><br>(Apellis)  | MN, IgAN, C3G,<br>Dense Deposit Disease,<br>or Lupus Nephritis | 18+          | APL-2                       | ≥ 30              | 2     |
| <b>A PHASE IB/2A STUDY OF VB119 IN PRIMARY MN</b><br>(ValenzaBio)  | MN   | 18+          | VB119                       | ≥ 45              | 1 & 2 |
| <b>NEW-PLACE (ENROLLMENT COMPLETE)</b><br>(MorphoSys)  | MN   | 18-80        | MOR202                      | > 30              | 2     |
| <b>MONET (ITALY)</b><br>(MorphoSys)  | MN   | 18+          | MOR202                      | > 30              | 2     |
| <b>M-PLACE (ENROLLMENT COMPLETE)</b><br>(MorphoSys)  | MN   | 18-80        | MOR202                      | ≥ 30              | 1 & 2 |
| <b>REBOOT</b><br>(NIAID)   | MN, Nephrotic<br>Syndrome                                      | 18-75        | Belimumab,<br>Rituximab     | ≥ 30              | 2     |
| <b>EFFICACY &amp; SAFETY OF OBINUTUZUMAB IN PARTICIPANTS WITH PRIMARY MN</b><br>(Hoffmann-La Roche)                              | MN   | 18-75        | Obinutuzumab                | ≥ 30              | 3     |
| <b>RENEW (ENROLLMENT COMPLETE)</b><br>(BioCryst)   | MN, IgAN, C3G  | 18+          | BCX9930                     | ≥ 50              | 2     |
| <b>STUDY OF ANG-3070 IN PRIMARY GLOMERULAR DISEASE</b><br>(Angion)   | Glomerular Disease   | 18+          | ANG-3070                    | ≥ 40              | 2     |
| <b>STUDY OF BI 690517 ALONE AND IN COMBINATION WITH EMPAGLIFLOZIN IN DIABETIC AND NON-DIABETIC CKD</b><br>(Boehringer Ingelheim) | Chronic Kidney Disease   | 18+          | BI 690517                   | ≥ 30 &<br>< 90    | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI 685509 ON KIDNEY FUNCTION IN CKD WITHOUT DIABETES</b><br>(Boehringer Ingelheim) | Chronic Kidney Disease   | 18+          | BI 685509                   | ≥ 20 &<br>< 90    | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)  | Chronic Kidney Disease   | 18+          | AZD5718,<br>Dapagliflozin   | 20-75             | 2     |
| <b>ZENITH-CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease   | 18+          | Zibotentan<br>Dapagliflozin | ≥ 20              | 2     |
| <b>EAGLE</b><br>(Reata)  | Chronic Kidney Disease,<br>Alport Syndrome                     | 12+          | Bardoxolone<br>methyl       | ≥ 20              | 3     |

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# CLINICAL TRIALS FOR MEMBRANOUS PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE         | DRUG/<br>COMPOUND                 | EGFR<br>IN ML/MIN | PHASE |
|--|--|----------------------|-----------------------------------|-------------------|-------|
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE WITH CKD</b><br>(Bayer)   | Chronic Kidney Disease   | 6 months to 17 years | Finerenone (Kerendia, BAY94-8862) | ≥ 30              | 3     |
| <b>FIND-CKD</b><br>(Bayer)   | Chronic Kidney Disease   | 18+                  | Finerenone (BAY94-8862)           | ≥ 25 and < 90     | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi)                              | Secondary Hyperparathyroidism<br>Chronic Kidney Disease        | 5-18                 | Doxercalciferol (GZ427397)        | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)   | Chronic Kidney Disease Stage IV                                | 18-80                | US-APR2020                        | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS AND SAFETY OF MK-2060 IN PATIENTS WITH ESRD RECEIVING HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)                               | End Stage Renal Disease & Kidney Failure                       | 18+                  | MK-2060                           | N/A               | 2     |
| <b>FREEDOM-1</b><br>(Talaris)  | De novo living donor transplant                                | 18+                  | FCR001                            | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)  | De novo living donor transplant (3-12 months after transplant) | 18+                  | FCR001                            | N/A               | 2     |
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant   | 18-75                | Extracorporeal photopheresis      | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor transplant                                | 18-65                | Apheresis                         | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant   | 18-70                | VIB4920 & Belatacept              | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor transplant                                | 18-70                | CD28 & IL-6 Receptor Antagonists  | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant rejection                                    | 18-70                | TX200-TR101                       | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR C3G PATIENTS

| TRIAL NAME<br>(sponsor)   | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE               | DRUG/<br>COMPOUND                       | EGFR<br>IN ML/MIN | PHASE |
|---|--|----------------------------|---|-------------------|-------|
| <b>APPEAR-C3G</b><br>(Novartis)   | C3G  | 18-60                      | LNP023                                  | ≥ 30              | 3     |
| <b>VALIANT</b><br>(Apellis)   | C3G, IC-MPGN, C3GN,<br>DDD, MPGN                               | 12+                        | Pegcetacoplan                           | ≥ 30              | 3     |
| <b>THE NOBLE STUDY</b><br>(Apellis)   | C3G, C3GN, DDD,<br>MPGN, or IC-MPGN<br>that has recurred after | 18+                        | Pegcetacoplan                           | ≥ 15              | 2     |
| <b>THE DISCOVERY TRIAL (ENROLLMENT COMPLETE)</b><br>(Apellis)   | C3G, Dense Deposit<br>Disease, IgAN, MN, or<br>Lupus Nephritis | 18+                        | APL-2                                   | ≥ 30              | 2     |
| <b>RENEW (ENROLLMENT COMPLETE)</b><br>(BioCryst)  | C3G, MN, IgAN  | 18+                        | BCX9930                                 | ≥ 50              | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI<br/>685509 ON KIDNEY FUNCTION IN CKD WITHOUT<br/>DIABETES</b><br>(Boehringer Ingelheim)                  | Chronic Kidney Disease   | 18+                        | BI 685509                               | ≥ 20 &<br>< 90    | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease   | 18+                        | AZD5718,<br>Dapagliflozin               | 20-75             | 2     |
| <b>ZENITH-CKD</b><br>(AstraZeneca)  | Chronic Kidney Disease   | 18+                        | Zibotentan<br>Dapagliflozin             | ≥ 20              | 2     |
| <b>EAGLE</b><br>(Reata)   | Chronic Kidney Disease,<br>Alport Syndrome                     | 12+                        | Bardoxolone<br>methyl                   | ≥ 20              | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI<br/>OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE<br/>WITH CKD</b><br>(Bayer)                | Chronic Kidney Disease   | 6 months<br>to 17<br>years | Finerenone<br>(Kerendia,<br>BAY94-8862) | ≥ 30              | 3     |
| <b>FIND-CKD</b><br>(Bayer)  | Chronic Kidney Disease   | 18+                        | Finerenone<br>(BAY94-8862)              | ≥ 25 and<br>< 90  | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD<br/>STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON<br/>DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi) | Secondary<br>Hyperparathyroidism<br>Chronic Kidney Disease     | 5-18                       | Doxercalciferol<br>(GZ427397)           | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)  | Chronic Kidney Disease<br>Stage IV                             | 18-80                      | US-APR2020                              | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS<br/>AND SAFETY OF MK-2060 IN PATIENTS WITH ESRD<br/>RECEIVING HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)  | End Stage Renal Disease<br>& Kidney Failure                    | 18+                        | MK-2060                                 | N/A               | 2     |

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# CLINICAL TRIALS FOR C3G PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE | DRUG/<br>COMPOUND                | EGFR<br>IN ML/MIN | PHASE |
|--|--|--------------|----------------------------------|-------------------|-------|
| <b>FREEDOM-1</b><br>(Talaris)  | De novo living donor transplant                                | 18+          | FCR001                           | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)  | De novo living donor transplant (3-12 months after transplant) | 18+          | FCR001                           | N/A               | 2     |
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)  | De novo transplant   | 18-75        | Extracorporeal photopheresis     | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor transplant                                | 18-65        | Apheresis                        | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio) | De novo transplant   | 18-70        | VIB4920 & Belatacept             | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b>         | De novo living donor transplant                                | 18-70        | CD28 & IL-6 Receptor Antagonists | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant rejection                                    | 18-70        | TX200-TR101                      | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR C3GN PATIENTS

| TRIAL NAME<br>(sponsor)   | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE            | DRUG/<br>COMPOUND                       | EGFR<br>IN ML/MIN | PHASE |
|---|--|-------------------------|---|-------------------|-------|
| <b>THE NOBLE STUDY</b><br>(Apellis)   | C3GN, C3G, DDD,<br>MPGN, or IC-MPGN<br>that has recurred after       | 18+                     | Pegcetacoplan                           | ≥ 15              | 2     |
| <b>VALIANT</b><br>(Apellis)   | C3GN, C3G, IC-MPGN,<br>DDD, MPGN                                     | 12+                     | Pegcetacoplan                           | ≥ 30              | 3     |
| <b>THE DISCOVERY TRIAL (ENROLLMENT COMPLETE)</b><br>(Apellis)   | C3GN, C3G, IC-MPGN,<br>DDD, MPGN                                     | 18+                     | Pegcetacoplan                           | ≥ 30              | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI 685509<br/>ON KIDNEY FUNCTION IN CKD WITHOUT DIABETES</b><br>(Boehringer Ingelheim)                      | Chronic Kidney Disease   | 18+                     | BI 685509                               | ≥ 20 &<br>< 90    | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease   | 18+                     | AZD5718,<br>Dapagliflozin               | 20-75             | 2     |
| <b>ZENITH-CKD</b><br>(AstraZeneca)  | Chronic Kidney Disease   | 18+                     | Zibotentan<br>Dapagliflozin             | ≥ 20              | 2     |
| <b>EAGLE</b><br>(Reata)   | Chronic Kidney Disease,<br>Alport Syndrome                           | 12+                     | Bardoxolone<br>methyl                   | ≥ 20              | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI<br/>OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE<br/>WITH CKD</b><br>(Bayer)                | Chronic Kidney Disease   | 6 months<br>to 17 years | Finerenone<br>(Kerendia,<br>BAY94-8862) | ≥ 30              | 3     |
| <b>FIND-CKD</b><br>(Bayer)  | Chronic Kidney Disease   | 18+                     | Finerenone<br>(BAY94-8862)              | ≥ 25 and<br>< 90  | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD<br/>STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON<br/>DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi) | Secondary<br>Hyperparathyroidism<br>Chronic Kidney Disease           | 5-18                    | Doxercalciferol<br>(GZ427397)           | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)  | Chronic Kidney Disease<br>Stage IV                                   | 18-80                   | US-APR2020                              | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS AND<br/>SAFETY OF MK-2060 IN PATIENTS WITH ESRD RECEIVING<br/>HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)  | End Stage Renal Disease<br>& Kidney Failure                          | 18+                     | MK-2060                                 | N/A               | 2     |
| <b>FREEDOM-1</b><br>(Talaris)   | De novo living donor<br>transplant                                   | 18+                     | FCR001                                  | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)   | De novo living donor<br>transplant (3-12 months<br>after transplant) | 18+                     | FCR001                                  | N/A               | 2     |

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# CLINICAL TRIALS FOR C3GN PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS            | AGE<br>RANGE | DRUG/<br>COMPOUND                | EGFR<br>IN ML/MIN | PHASE |
|--|---------------------------------|--------------|----------------------------------|-------------------|-------|
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant              | 18-75        | Extracorporeal photopheresis     | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor transplant | 18-65        | Apheresis                        | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant              | 18-70        | VIB4920 & Belatacept             | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor transplant | 18-70        | CD28 & IL-6 Receptor Antagonists | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant rejection     | 18-70        | TX200-TR101                      | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR MPGN PATIENTS

| TRIAL NAME<br>(sponsor)   | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE | DRUG/<br>COMPOUND                 | EGFR<br>IN ML/MIN | PHASE |
|---|--|--------------|-----------------------------------|-------------------|-------|
| <b>THE NOBLE STUDY</b><br>(Apellis)   | MPGN, IC-MPGN, C3GN, C3G, or DDD, or any of these conditions that has recurred after kidney transplant | 18+          | Pegcetacoplan                     | ≥ 15              | 2     |
| <b>VALIANT</b><br>(Apellis)   | MPGN, IC-MPGN, C3G, C3GN, DDD  | 12+          | Pegcetacoplan                     | ≥ 30              | 3     |
| <b>STUDY OF ANG-3070 IN PRIMARY GLOMERULAR DISEASE</b><br>(Angion)  | Glomerular Disease   | 18+          | ANG-3070                          | ≥ 40              | 2     |
| <b>STUDY TO TEST EFFECT OF DIFFERENT DOSES OF BI 685509 ON KIDNEY FUNCTION IN CKD WITHOUT DIABETES</b><br>(Boehringer Ingelheim)                  | Chronic Kidney Disease   | 18+          | BI 685509                         | ≥ 20 & < 90       | 2     |
| <b>STUDY OF AZD5718 IN PROTEINURIC CKD</b><br>(AstraZeneca)   | Chronic Kidney Disease   | 18+          | AZD5718, Dapagliflozin            | 20-75             | 2     |
| <b>ZENITH-CKD (ENROLLMENT COMPLETE)</b><br>(AstraZeneca)  | Chronic Kidney Disease   | 18+          | Zibotentan Dapagliflozin          | ≥ 20              | 2     |
| <b>EAGLE</b><br>(Reata)   | Chronic Kidney Disease, Alport Syndrome  | 12+          | Bardoxolone methyl                | ≥ 20              | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE WITH CKD</b><br>(Bayer)                | Chronic Kidney Disease   | 6 to 17      | Finerenone (Kerendia, BAY94-8862) | ≥ 30              | 3     |
| <b>STUDY OF FINERENONE IN NON-DIABETIC CHRONIC KIDNEY DISEASE</b><br>(Bayer)  | Chronic Kidney Disease   | 18+          | Finerenone (BAY94-8862)           | ≥ 25 and < 90     | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi) | Secondary Hyperparathyroidism<br>Chronic Kidney Disease  | 5-18         | Doxercalciferol (GZ427397)        | 15-59             | 3     |
| <b>SAFETY AND EFFICACY OF APR2020 IN CKD STAGE IV</b><br>(Kibow)  | Chronic Kidney Disease Stage IV  | 18-80        | US-APR2020                        | 15-29             | 2     |
| <b>PREVENTION OF ARTERIOVENOUS GRAFT THROMBOSIS AND SAFETY OF MK-2060 IN PATIENTS WITH ESRD RECEIVING HEMODIALYSIS</b><br>(Merck Sharpe & Dohme)  | End Stage Renal Disease & Kidney Failure   | 18+          | MK-2060                           | N/A               | 2     |
| <b>FREEDOM-1</b><br>(Talaris)   | De novo living donor transplant  | 18+          | FCR001                            | N/A               | 3     |
| <b>FREEDOM-2</b><br>(Talaris)   | De novo living donor transplant (3-12 months after transplant)   | 18+          | FCR001                            | N/A               | 2     |

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# CLINICAL TRIALS FOR MPGN PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS            | AGE<br>RANGE | DRUG/<br>COMPOUND                | EGFR<br>IN ML/MIN | PHASE |
|--|---------------------------------|--------------|----------------------------------|-------------------|-------|
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant              | 18-75        | Extracorporeal photopheresis     | N/A               | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor transplant | 18-65        | Apheresis                        | N/A               | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant              | 18-70        | VIB4920 & Belatacept             | N/A               | 2     |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor transplant | 18-70        | CD28 & IL-6 Receptor Antagonists | N/A               | 1 & 2 |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant rejection     | 18-70        | TX200-TR101                      | N/A               | 1 & 2 |

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# CLINICAL TRIALS FOR TRANSPLANT PATIENTS

| TRIAL NAME<br>(sponsor)  | PATIENT<br>DIAGNOSIS   | AGE<br>RANGE | DRUG/<br>COMPOUND                | EGFR<br>IN ML/MIN                  | PHASE |
|--|--|--------------|----------------------------------|------------------------------------|-------|
| <b>FREEDOM-1</b><br>(Talaris)  | De novo living donor transplant                                | 18+          | FCR001                           | N/A                                | 3     |
| <b>FREEDOM-2</b><br>(Talaris)  | De novo living donor transplant (3-12 months after transplant) | 18+          | FCR001                           | N/A                                | 2     |
| <b>PHOTOPHERESIS IN THE PREVENTION OF ACUTE KIDNEY REJECTION IN HIGHLY SENSITIZED DE NOVO TRANSPLANT RECIPIENTS (SPAIN - COMING SOON)</b><br>(Mallinckrodt)                    | De novo transplant   | 18-75        | Extracorporeal photopheresis     | N/A                                | N/A   |
| <b>AUTOLOGOUS REGULATORY T CELL INFUSION AND ZORTRESS IN RENAL TRANSPLANT RECIPIENTS</b><br>(Novartis)   | De novo living donor transplant                                | 18-65        | Apheresis                        | N/A                                | N/A   |
| <b>STUDY OF VIB4920 AND BELATACEPT FOR PROPHYLAXIS OF ALLOGRAFT REJECTION IN KIDNEY TRANSPLANT (ENROLLMENT COMPLETE)</b><br>(Horizon Therapeutics/Viela Bio)                   | De novo transplant   | 18-70        | VIB4920 & Belatacept             | N/A                                | 2     |
| <b>THE NOBLE STUDY</b><br>(Apellis)  | MPGN, IC-MPGN, C3GN, C3G, or DDD, or any of these conditions   | 18+          | Pegcetacoplan                    | ≥ 15                               | 3     |
| <b>VALIANT</b><br>(Apellis)  | MPGN, IC-MPGN, C3G, C3GN, DDD                                  | 12+          | Pegcetacoplan                    | ≥ 30                               | N/A   |
| <b>POST-APPROVAL STUDY OF LIPOSORBER LA-15 SYSTEM FOR PEDIATRIC &amp; DRUG RESISTANT ADULT PATIENTS WITH FSGS</b><br>(Kaneka)  | Post-transplant recurrent FSGS or Primary FSGS                 | 5-75         | Liposorber LA-15 System          | ≥ 45 or post-transplant recurrence | 1 & 2 |
| <b>T CELL MODULATION IN KIDNEY TRANSPLANTATION WITH BIOLOGIC BLOCKADE OF DUAL EFFECTOR PATHWAYS, CD28 &amp; IL-6 (CTOT-24) (ENROLLMENT COMPLETE)</b><br>(Bristol-Myers Squibb) | De novo living donor transplant                                | 18-70        | CD28 & IL-6 Receptor Antagonists | 1 & 2                              | 2     |
| <b>STEADFAST</b><br>(Sangamo Therapeutics)   | Kidney transplant rejection                                    | 18-70        | TX200-TR101                      | 1 & 2                              |       |

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# CLINICAL TRIALS FOR PEDIATRIC PATIENTS

| TRIAL NAME<br>(sponsor)   | PATIENT<br>DIAGNOSIS                                       | AGE<br>RANGE            | DRUG/<br>COMPOUND                       | EGFR<br>IN ML/MIN                         | PHASE |
|---|--|-------------------------|---|---|-------|
| <b>EPPIK</b><br>(Traverse)  | FSGS, MCD, IgAN,<br>IgAV, Alport Syndrome                  | 1-17                    | Sparsentan                              | ≥ 30                                      | 2     |
| <b>DUPLEX (ENROLLMENT COMPLETE)</b><br>(Traverse)   | FSGS   | 8-75                    | Sparsentan                              | ≥ 30                                      | 3     |
| <b>VALIANT</b><br>(Apellis)   | C3G, IC-MPGN, C3GN,<br>DDD, MPGN                           | 12+                     | Pegcetacoplan                           | ≥ 30                                      | 3     |
| <b>POST-APPROVAL STUDY OF LIPOSORBER LA-15 SYSTEM FOR PEDIATRIC &amp; DRUG RESISTANT ADULT PATIENTS WITH FSGS</b><br>(Kaneka)                     | Post-transplant recurrent<br>FSGS or Primary FSGS          | 5-75                    | Liposorber LA-15<br>System              | ≥ 45 or post-<br>transplant<br>recurrence | N/A   |
| <b>TUMOR NECROSIS FACTOR INHIBITION IN FSGS AND TREATMENT-RESISTANT MCD</b><br>(University of Michigan)   | FSGS, MCD  | 6-70                    | adalimumab                              | > 45                                      | 2     |
| <b>EAGLE</b><br>(Reata)   | Chronic Kidney Disease,<br>Alport Syndrome                 | 12+                     | Bardoxolone<br>methyl                   | ≥ 20                                      | 3     |
| <b>FIONA - STUDY OF FINERENONE IN ADDITION TO AN ACEI OR ARB FOR CHILDREN 6 MONTHS TO &lt; 18 YEARS OF AGE WITH CKD</b><br>(Bayer)                | Chronic Kidney Disease                                     | 6 months to<br>17 years | Finerenone<br>(Kerendia,<br>BAY94-8862) | ≥ 30                                      | 3     |
| <b>STUDY OF HECTOROL IN PEDIATRIC PATIENTS WITH CKD STAGES 3 AND 4 WITH HYPERPARATHYROIDISM NOT ON DIALYSIS (ENROLLMENT COMPLETE)</b><br>(Sanofi) | Secondary<br>Hyperparathyroidism<br>Chronic Kidney Disease | 5-18                    | Doxercalciferol<br>(GZ427397)           | 15-59                                     | 3     |

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