

## FIND Medical Record Review

Barcode and  
Participant ID

Study Coordinator ID \_\_\_\_\_

Date of review (MM/DD/YYYY) \_\_\_\_ | \_\_\_\_ | \_\_\_\_

**Instruction to record abstractor:** During review of the medical record(s), please review the Medical Questionnaire (Form 01), since it may be necessary to add or correct some data on the questionnaire (e.g., medicines or date of onset of diabetes or dialysis).

### RECORD TYPE

1. Indicate type of record(s) reviewed (*Mark all that apply*).

- Dialysis unit
- Primary care (*Including records from the endocrinologist and/or nephrologist*)
- Hospital
- Other \_\_\_\_\_

## PARTICIPANT INFORMATION

2. Name: \_\_\_\_\_  
*Last First Middle Maiden*

3. Birth date (MM/DD/YYYY): \_\_\_\_ | \_\_\_\_ | \_\_\_\_

4. Sex: Male Female

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## KIDNEY DISEASE

### Kidney Disease

5. Is the level of urinary protein/albumin recorded in the record?

No  Yes

If **Yes**, then specify the following (*Mark the maximum value only*):

	<u>Protein excretion rate</u>		<u>P/C ratio</u>		<u>Albumin excretion rate</u>		<u>A/C ratio</u>
a. <input type="checkbox"/>	<50 mg/24h	or	<0.15 mg/mg	or	<30 mg/24h	or	<0.03 mg/mg
b. <input type="checkbox"/>	≥50 mg/24h	or	≥0.15 mg/mg	or	≥30 mg/24h	or	≥0.03 mg/mg
c. <input type="checkbox"/>	≥500 mg/24h	or	≥0.5 mg/mg	or	≥300 mg/24h	or	≥0.3 mg/mg
d. <input type="checkbox"/>	≥1.0 g/24h	or	≥1.0 mg/mg	or	≥1.0 g/24h	or	≥1.0 mg/mg
e. <input type="checkbox"/>	≥3.0 g/24h	or	≥3.0 mg/mg	or	≥3.0 g/24h	or	≥3.0 mg/mg
f. <input type="checkbox"/>	Nephrotic ( 3.0g – 3.5g)						
g. <input type="checkbox"/>	None						

If **a**, record date of **LAST** value (*MM/DD/YYYY*): \_\_\_ | \_\_\_ | \_\_\_\_\_

Also record whether subject is receiving antihypertensive therapy at **LAST** value:  No  Yes  
**HIGHEST** value:  No  Yes

If **e**, record date of **FIRST** value at this level (*MM/DD/YYYY*): \_\_\_ | \_\_\_ | \_\_\_\_\_

#### ***Equivalent Measures of Urinary Albumin Excretion***

30 mg/24h	=	20 µg/min
300 mg/24h	=	200 µg/min

If the excretion reported above is a ratio, record the urine protein (albumin) and creatinine concentrations:

Urine protein (*mg/l*) \_\_\_\_\_ . \_\_\_\_ or Urine albumin (*mg/l*) \_\_\_\_\_ . \_\_\_\_

Urine creatinine (*g/l*) \_\_\_\_ . \_\_\_\_

6. Is the participant receiving chronic renal replacement therapy?

No  Yes If **Yes**, record the date of onset (*MM/DD/YYYY*): \_\_\_ | \_\_\_ | \_\_\_\_\_  
 Yes AA MALD with Nephropathy but no DM

If **No**, record the **HIGHEST** serum creatinine concentration (*mg/dl*): \_\_\_ . \_\_\_\_

Date of **HIGHEST** serum creatinine concentration (*MM/DD/YYYY*): \_\_\_ | \_\_\_ | \_\_\_\_\_

7. Primary cause of kidney failure / renal insufficiency (*Mark all that apply*)?

- |  |  |
|--|--|
| <input type="checkbox"/> Diabetes                      | <input type="checkbox"/> Polycystic kidney disease |
| <input type="checkbox"/> Hypertension                  | <input type="checkbox"/> Lupus nephritis           |
| <input type="checkbox"/> IgA nephropathy               | <input type="checkbox"/> Kidney cancer             |
| <input type="checkbox"/> Membranous glomerulonephritis | <input type="checkbox"/> Obstruction               |
| <input type="checkbox"/> Focal glomerulosclerosis      | <input type="checkbox"/> Don't know                |
| <input type="checkbox"/> Other _____                   | <input type="checkbox"/> None                      |

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8. Is a kidney biopsy recorded in the record?

No  Yes If **Yes**, record the biopsy date (MM/DD/YYYY): \_\_\_ | \_\_\_ | \_\_\_\_\_

If **Yes**, specify methods of evaluation (Mark all that apply).

Light microscopy  Electron microscopy  Immunofluorescence

Also specify the histologic findings (Mark all that apply).

- Increased nodular mesangial matrix.
- Increased diffuse mesangial matrix.
- Thickened glomerular basement membrane.
- Arterial hyalinization.
- Arteriolar hyalinization.
- Mesangial immunoglobulin or paraprotein deposits by immunofluorescence.
- Amyloid deposits by Congo red staining or electron microscopy.
- Electron dense deposits within the glomerular basement membrane or glomerular capillary subendothelial space.
- Non-Diabetic Pathological Diagnosis

HISTORY OF DIABETES

9. Has a diagnosis of diabetes been made?

No  Yes

If **Yes**, record the source of the diagnosis (Mark all that apply):

- Fasting plasma glucose  $\geq 126$  mg/dl (or venous whole blood glucose  $\geq 110$  mg/dl or capillary whole blood glucose  $\geq 110$  mg/dl) \*
- Random plasma glucose  $\geq 200$  mg/dl (or venous whole blood glucose  $\geq 180$  mg/dl or capillary whole blood glucose  $\geq 200$  mg/dl) \* and symptoms (*polyuria, polydipsia, polyphagia*).
- Two-hour post-load plasma glucose  $\geq 200$  mg/dl (or venous whole blood glucose  $\geq 180$  mg/dl or capillary whole blood glucose  $\geq 200$  mg/dl) \* (OGTT).
- Clinical diagnosis without documented plasma glucose concentration.

Record the earliest date of diagnosis (MM/DD/YYYY): \_\_\_ | \_\_\_ | \_\_\_\_\_

\*The measurement of glucose in serum is discouraged by the WHO (World Health Organization). Unless the red cells are immediately removed to prevent glycolysis, serum samples should not be used for diagnosing diabetes. If only serum glucose values are found in the record, however, they should be interpreted as if they were plasma values.

10. Is a diagnosis of diabetic ketoacidosis or diabetic coma recorded in the record?

No  Yes

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## DIABETIC EYE DISEASE

11. Is a diagnosis of diabetic retinopathy recorded in the record?

No       Yes

If **Yes**, specify the severity of the retinopathy (*Mark all that apply*).

Background retinopathy

Pre-proliferative retinopathy

Proliferative retinopathy

Vitreous hemorrhage

Macular edema

Photocoagulation therapy