

## Global strategies

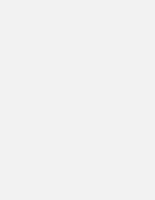
• WHO has been outlining strategies to reduce mortality

from chronic non-communicable diseases by 2022.

• CDC have launched public health strategies to prevent the

development, progression and complications of the disease

in the United States.





# CKD Definition

- Kidney damage for  $\geq$ 3 months,
  - **as defined by:**
- structural or functional abnormalities of the kidney with or without decreased glomerular filtration rate (GFR) manifested by one or more of the following features:
  - abnormalities in the composition of the blood or urine,
  - abnormalities in imaging tests,
  - abnormalities on kidney biopsy;



# CKD Definition

- GFR <60 ml/min/1.73 m<sup>2</sup> for  $\ge$ 3 months
  - with or without the other above-mentioned signs of kidney damage.

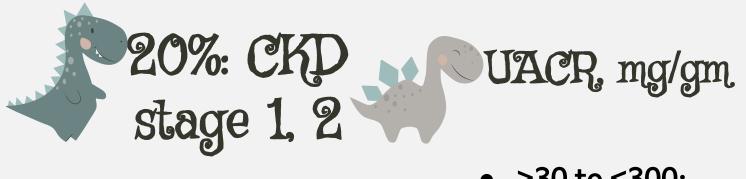




#### An Analysis of the CURE-CKD Registry

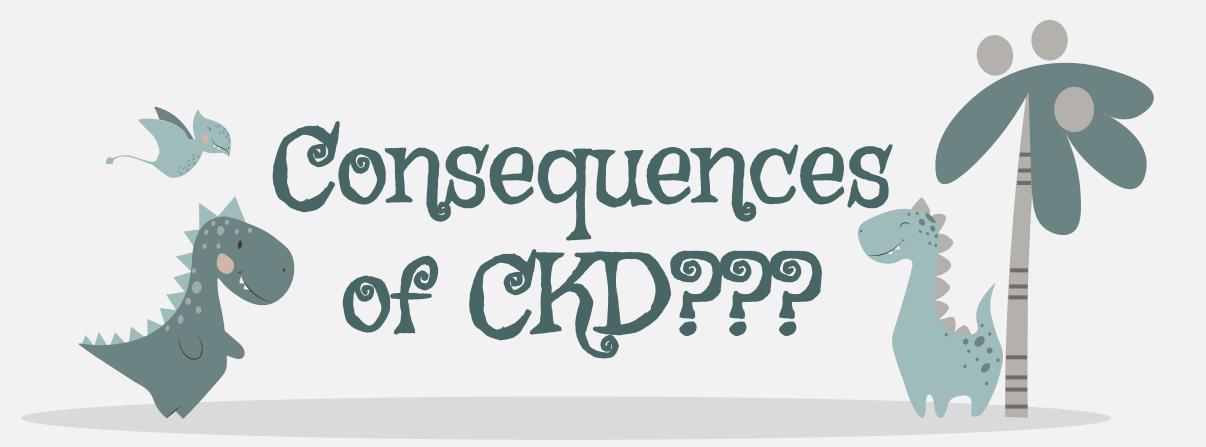
prevalence & demographic- data





 Non-Latino white: 52.8% • 43.8% boys

- >30 to ≤300:
  - 1.6%



This is where you section ends. Duplicate this set of slides as many times you need to go over all your sections.

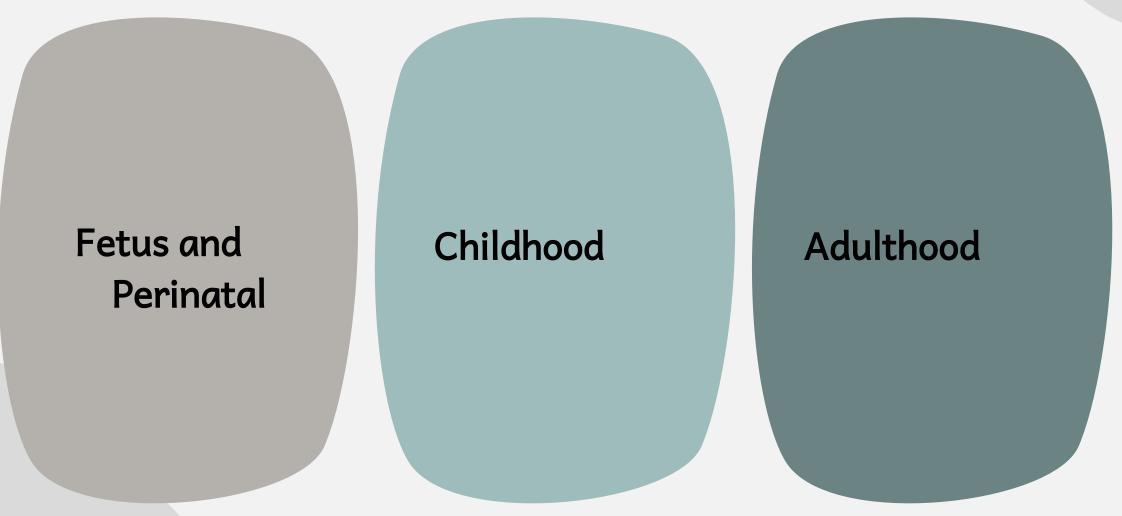
#### Cardiovascular Disease risk factors

- Hypercalcemia,
- Hyperphosphatemia,
- High parathyroid hormone (PTH),
- Very low or high vitamin D levels
- Anemia
- Malnutrition

- A multitude of middle and large molecular weight substances together labelled 'uremic toxins',
- Chronic inflammation,
- Oxidative stress,
- Hyper-homocysteinemia



#### Levels of Prevention

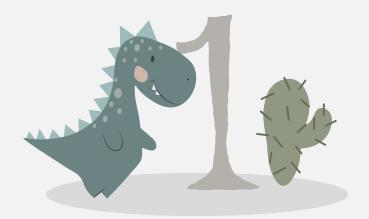




## Did you know?

- In 1986, the epidemiologist **David Barker** proposed the theory of a fetal and infantile origin for adult diseases:
  - "Adverse environments during fetal life and early childhood imply an increased risk of illness during adulthood."





#### Fetal programming and chronic adult diseases

- In Pediatrics, kidney diseases can be "silent" with nonspecific signs and symptoms.
- The first responsible doctors are pediatricians and family doctors.
- In fact, prevention of kidney disease must begin before a woman becomes pregnant.
  - It is called fetal and infant programming of noncommunicable chronic diseases of adults (NCDA).

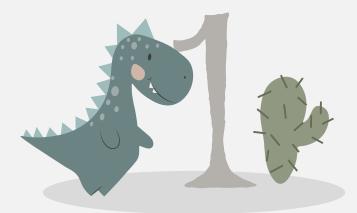
## Fetal Programming

Unfavorable intrauterine environment → inhospitable experiences suffered by the embryo/fetus → adaptations of the embryo/fetus (epigenetic phenomena) → impaired renal maturation "FETAL AND PERINATAL PROGRAMMING OF KIDNEY DISEASES IN ADULTS"



Help Children to save future Adults

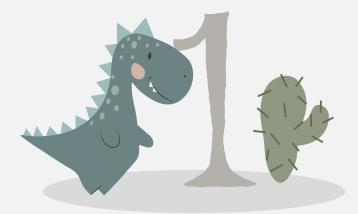




## Factors induce fetal

#### programming:

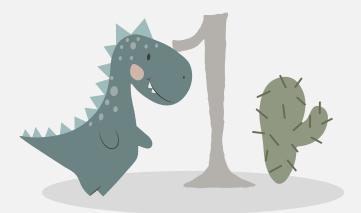
- I. Maternal:
- Maternal nutrition (mainly proteins and glucose);
- Consumption of illicit drugs alcohol and tobacco
- Hypertension;
- Pre-eclampsia;
- Primiparity;
- Maternal overweight / obesity;
- Mother born small for gestational age or with fetal growth restriction;
- Asthma;
- Polycystic ovary syndrome.



Factors induce fetal programming:

- II. Placental:
- Changes in uterus placental circulation;
- Changes in nutrient transfer;
- Placental infarction;
- Abnormal development of the placenta.





# Factors induce fetal programming:

- III. Fetal:
- Chromosomal abnormalities.

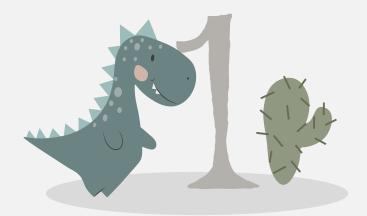
Regarding kidney diseases, unfavorable intrauterine environment could cause adaptations of the embryo/fetus resulting in compromised renal maturation. This sequence of events is called *fetal and perinatal programming of kidney diseases in adults*.



#### Prevention of kidney diseases in childhood

 Considering all the evidence, in April 2016 a Workshop was held with the aim of highlighting the association between fetal and child development and the increased risk of adult diseases, with a focus on arterial hypertension and CKD.



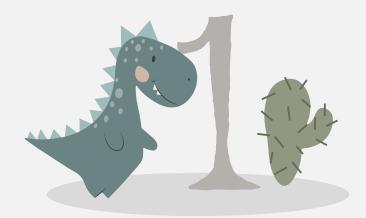


# Identify Children at risk for CKD.

#### • who should be screened?

- I. Family history of CKD or other genetic kidney disease
- II. Family history of hypertension, diabetes and cardiovascular disease in parents, uncles and grandparents
- III. Low birth weight and premature babies

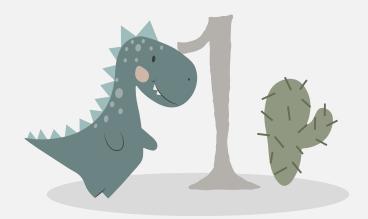




#### Identify Children at risk for CKD continue

- IV. History of long hospital stay in the neonatal period
- V. Renal dysplasia / hypoplasia
- VI. Spinal cord tumors and traumas
- VII. Congenital malformations of the urinary tract





#### Identify Children at risk for CKD continue

- VIII. Previous history of hemolytic-uremic syndrome
- IX. Previous history of glomerulopathies
- X. Overweight / obese children
- XI. Bladder diseases (neurogenic, non-neurogenic bladder, lower urinary tract dysfunction).



### Different aspects of CKD Prevention?

- Primary prevention aims to eliminate or reduce exposure to factors that cause renal disease.
- Secondary prevention in which the prevention of the progression of renal damage from stage 1 to stage 5 is carried out by introducing appropriate measures at various stages of CKD.
- Tertiary prevention strategies are focused on the: reduction or delay of long-term complications, impairments or disabilities in established disease, requiring renal replacement therapy (RRT).

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# Primary preventive strategies

- 1. Control of possible future pregnancy:
  - Avoid using drugs (ACEi, AG ll RBs, NSAIDs, illicit drugs)
- I. Control of overweight/obesity and metabolic syndrome
- II. Control of dyslipidemia
- III. Pay attention to the future mother's nutrition
- IV. Prevent disease (rubella, toxoplasmosis, cytomegalovirus, etc.)
- V. Guide on the importance of genetic counseling
- VI. Use folic acid.





#### 2. Control of the pregnant woman:

- Avoid using drugs (ACEi, AG ll RBs, NSAIDs, illicit drugs)
- I. Control of overweight / obesity and metabolic syndrome
- II. Control of dyslipidemia
- III. Pay attention to the future mother's nutrition
- IV. Prevent disease (rubella, toxoplasmosis, cytomegalovirus, etc.)
- V. Prohibit the use of tobacco and alcohol
- VI. Prevent prematurity, if possible
- VII. Early detection of intrauterine growth restriction.





#### 3. Infant control:

- Avoid using drugs (ACEi, AG ll RBs, NSAIDs, illicit drugs)
- I. Stimulate breastfeeding
- II. Take care of infant nutrition
- III. Attention to secondhand smoke
- IV. Watch out for postnatal catch up.





- 4. Child and adolescent control :
- I. Prevent overweight / obesity and dyslipidemia
- II. Promote food education
- III. Encourage physical activity
- IV. Encourage breastfeeding
- V. Prohibit the use of tobacco and alcohol.



## Primary preventive strategies

#### Parents and caregivers must have a healthy lifestyle:

- I. To avoid soft drinks, industrialized juices, sweets, sausages, canned goods, salt;
- II. Provide a diet rich in fruits, vegetables, milk and with a reduced content of saturated fats;
- III. Ensure adequate intake of potassium with many fruits;
- **IV**. To avoid quick and processed snacks;
- V. Do regular physical exercises and be an example;
- VI. To avoid second hand smoke;
- VII. To read labels to check the content of the food.



### Secondary preventive strategies

- 1. Postnatal preventive measures should include (pay attention to):
- I. Low birth weight and extremely premature infants
- II. Postnatal nutrition (Obesity, FTT)
- III. Monitoring of albuminuria and blood pressure
- IV. Appropriate and early approach to sepsis and the possibility of acute kidney injury
- V. Early correction of metabolic disorders (polyuria, hyperkalemia, bicarbonaturia, etc).
- VI. Beware of clinical situations where there is a reduction in the number of nephrons (renal agenesis, VUR IV and V, UPJO & UVJO



### Secondary preventive strategies

- VII. Attention to nephrotoxic drugs (ACEi, ARBs NSAIDs, antibiotics aminoglycosides)
- VIII. Be careful with the use of nephrotoxic contrasts
- IX. Early correction of hypovolemia and shock with rapid replacement and volume maintenance
- X. Appropriate and early approach to UTI
- XI. Appropriate and early approach to glomerulopathies, (proteinuria, BP, dyslipidemia)
- XII. Control of metabolic diseases (glycemic control, dyslipidemia, proteinuria, blood pressure)
- XIII. Attention to rare diseases.



# Tertiary preventive strategies

- I. Blood pressure control
- II. Proteinuria control
- III. Dyslipidemia control
- IV. Diet control (protein and phosphorus)
- V. Anemia control
- VI. Vitamin D normalization
- VII. Bone mineral disease control
- VIII. Metabolic acidosis control.



# "CKD is common, harmful and treatable"

– Irene M. Pepperberg



### References

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#### Doyouhave any questions?