March 6, 2026

08:30–09:00
Participant Registration

09:00–09:30
Institutional Welcome Addresses

09:30–10:30

Primary Adrenal Insufficiency: Diagnosis and Therapeutic Management

Chairs: Roberta Giordano (Turin), Mariacarolina Salerno (Naples)

- Primary adrenal insufficiency: present and future
- Clinical case: adrenal insufficiency in pediatric age
- Clinical case: adrenal insufficiency in adulthood

  Discussion

10:30–11:00 | Lecture

Transition care for adolescents and young adults with congenital adrenal hyperplasia

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11:00–11:30
Coffee Break

11:30–12:30

Chronic Hypoparathyroidism

- Good clinical practice recommendations for the diagnosis and management of hypoparathyroidism
- Clinical case: hypoparathyroidism in pediatric age
- Clinical case: hypoparathyroidism in adulthood

Discussion

12:30–13:40

Male Hypogonadism

- Prenatal diagnosis and pediatric management of Klinefelter syndrome
- Pubertal induction in male patients with hypogonadism
- Male fertility preservation

Discussion

13:40–14:45
Light Lunch

14:45–15:50

Management of Prader–Willi Syndrome from Childhood to Adulthood

- Recommendations for the use of rhGH
- Strategies for obesity management
- Clinical case: transition of the patient with Prader–Willi syndrome
  Discussion

15:50–16:45

New Perspectives in Neonatal Diabetes and Acquired Hypothalamic Obesity

- Neonatal and pediatric diabetes: therapeutic perspectives between new formulations and cell transplantation
- Acquired hypothalamic obesity

Discussion

16:45–17:15
Coffee Break

17:15–18:30

Round Table

Unmet Needs: Dialogue Between Patient Associations and Clinicians

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March 7, 2026

09:00–10:00

Overview of Therapy for Acromegaly and Gigantism

- Acromegaly: indications for different therapeutic strategies
- Clinical case: pediatric gigantism
- Clinical case: adult acromegaly

Discussion

10:00–10:30 | Lecture

Outcomes in patients treated with rhIGF-1 for severe growth failure: real-world IGFD Registry data

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10:30–11:00
Coffee Break

11:00–12:15

Rare Thyroid Diseases

- Advances in diagnosis and treatment of medullary thyroid carcinoma
- Genetic disorders of thyroid hormone transport (MCT8 mutations), metabolism, and action (RTHα and RTHβ)
- Clinical case: central hypothyroidism

Discussion

12:15–12:45 | Lecture

The power of registries for rare endocrine diseases

12:45–13:00
Concluding Remarks