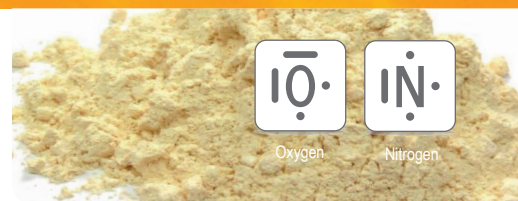


O, N determination in CeO₂ samples



Suitable analyzers

- ON 900
- ONH 2000

Used accessories

- Graphite crucibles 90190 or 90185 and 90180
- Suitable calibration material (NIST or other)
- Nickel capsule (90257)

Settings

- | | |
|-------------------------------|---------------------|
| ■ Comparator level: 20 mV | ■ Time |
| ■ Minimum time: 15 sec | Purge: 15 sec |
| ■ Maximum time: 1:30 min | Stability: 30 sec |
| ■ Post waiting: 10 sec | ■ Integration delay |
| ■ Base line deviation: 20 mV; | IR cell: 2 sec |
| Step: 16 mV; Time: 15 sec | TC cell: 10 sec |
| ■ Mode: continuous | ■ Analysis (1) |
| ■ Outgas | Time: 60 sec |
| Time: 45 sec | Power (1): 4.75 kW |
| Power: 6.0 kW | Power (2): 4.75 kW |


 Impulse
furnace

Sample preparation

Make sure that the CeO₂ sample is dry, otherwise remove the moisture at 105 °C. The dried sample is filled in the nickel capsule, and the nickel capsule has to be sealed completely to remove residual air.

Procedure

- Prepare ELTRA analyzer (exchange anhydron, sodium hydroxide, copper oxide or Schuetze reagent when necessary), clean furnace, sample drop mechanism, electrode tip
- Run three blanks with empty crucibles
- Calibrate the analyzer with suitable calibration material (NIST or other)
 - (1) Place empty crucible on the electrode tip, close furnace (F2 Button)
 - (2) Weigh calibration material (usually pins)
 - (3) Place calibration material in the sample drop mechanism and start analysis (F5 Button)

Repeat steps (1) – (3) at least three times;
Mark the results and use the calibration function in the software.

-> Now start with the actual analysis.

Typical results

Customer sample: CeO₂

Weight (mg)	% O	ppm N
15.63	18.57	99.0
15.76	18.48	91.7
15.47	18.38	101.4
15.05	18.49	114.2
15.68	18.63	107.9
15.62	18.60	97.5
15.96	18.30	105.6
15.44	18.33	109.7
15.24	18.12	100.8
15.32	18.52	106.9
Average values		
	18.44	103.5
Deviation		
	0.15 (0.85%)	6.6 (6.4%)