

MOERC (Metal Oxide Electrolysis Refinement Chamber)  
Sand → Silicon (200kg target) | 120kV + Battery | <\$10k v0.1

Earth → first, ISRU → ready benchtop line to convert ~1kg of oxide feed (beach sand, crushed rock, or simulant) into ~200kg of refined silicon using molten oxide electrolysis (MOE) followed by vacuum refining and ingot casting. No external chemical reductants (no carbon, no chlorosilanes). Designed around off-the-shelf parts and minimal fabrication.

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0) At a glance

Batch size: ~1kg dry feed (SiO<sub>2</sub> → dominant) → ~200kg Si (learning target; improved yields with practice)

Core process: MOE (Fe → Si → Al staging) → vacuum refining (P removal) → directional solidification

Operating temps: 1600 → 1700°C MOE cell; 1600 → 1900°C vacuum refine; 1450 → 1550°C casting

Power: 120kV AC, 15 → 20A; battery/inverter (~1.8kW) inline for ride-through; optional solar recharge

Atmosphere: low-leak sealed hot-zone; argon backfill (closed-loop reclaim); O<sub>2</sub> vent/capture

Footprint: bench stack, approx 0.8m × 0.6m × 1.2m total for all modules

Safety: high-T, high-brightness, O<sub>2</sub> evolution, off-gas management; multiple interlocks & E-stop

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1) System Architecture (modules)

Feed Prep & Autofeed

Hopper + auger feeder (vacuum-tolerant), inline dryer (150 → 250°C), screen (~250µm target), optional jaw crusher for rocks.

MOE Cell (Melter + Electrolyzer)

Insulated hot-zone, crucible/liner set, refractory metal electrodes, 1600 → 1700°C setpoint, staged potential for Fe → Si → Al.

Phase Separation

Weir/sump geometry to isolate Fe first, then collect molten Si away from slag/Fe (avoid Fe-Si alloying).

Vacuum Refining Pot

Dedicated vessel; 1600 → 1900°C at ~10<sup>-4</sup>Pa; cold-traps on exhaust to capture SiO and P species.

Ingot Casting

Directional solidification (Si → N<sub>2</sub> → coated crucibles) for multicrystalline bricks; optional small float-zone head for rods.

Gas & Vacuum Skid

Argon loop (buffer tank, dryer, filter), O<sub>2</sub> vent/capture, vacuum pump (HF-free process), baffles/cold-traps.

Controls & Sensing

PID/PLC, two-color pyrometry, LIBS (pre-feed/melt fingerprint), OES (process), RGA (vac/off-gas), interlocks.

## Power & Energy

120V mains → inverter/UPS (≈1.8kW continuous) → heater supplies; DC bus for controls; optional PV input.

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## 2) Bill of Materials (indicative, USD)

Prices are ballparks (new/COTS). Substitute equivalents as available.

### 2.1 Hot Zone & MOE Cell (~\$3,150)

High temp insulation kit (alumina fiber boards/blanket, 1700°C class), panels + fasteners → \$350

Modular steel frame (80/20 or welded), panels, hardware → \$250

Crucibles/liners: primary ZrO<sub>2</sub> liner (1.5" ID), backup Al<sub>2</sub>O<sub>3</sub> liners (x2) → \$400

Refractory metal electrodes: W cathode set + Mo spares; IrO<sub>2</sub> coated inert anode option (or consumable C anode minimized) → \$900

Induction/element heater set: high temp MoSi<sub>2</sub> elements (1700°C rating) w/ holders OR compact induction coil + 3" 5kW driver (120V input PFC) → \$950

Thermography: dual two color pyrometers (melt & wall), 600–2000°C → \$300

Fixtures: ceramic feed port, viewports (UV grade fused silica), fast shut baffles → \$150

### 2.2 Vacuum Refining Vessel (~\$1,750)

Small high vac furnace can (stainless shell, water cooled feedthroughs) → \$600

Heater: MoSi<sub>2</sub> element set or induction coil insert → \$450

Cold trap assembly (water cooled finger + removable cryo cup) → \$250

Vacuum hardware: KF25/KF40 set, valves, gauge (Pirani + capacitance manometer) → \$300

Si crucible (graphite or high density SiC not in contact with silica stage) → \$150

### 2.3 Gas & Vacuum (~\$1,600)

Rotary vane pump (5" 8m<sup>3</sup>/h) + backing filter → \$700

Dry scroll (used or small new) optional upgrade → \$700

Argon loop: 20–40L buffer tank, 0.1µm filter, desiccant dryer → \$200

### 2.4 Feed & Casting (~\$900)

Hopper + auger (stainless screw, NEMA 17/23 stepper, driver) → \$300

Inline dryer (500W cartridge + PID, 150–250°C) → \$150

Screens (250µm, 100µm), small jaw crusher (benchtop) → \$250

DS casting station: Si-Ni coated quartz crucibles (2–3), mold base & slow cool plate → \$200

### 2.5 Controls & Sensing (~\$1,900)

PLC or industrial micro (e.g., Codesys/Compact PLC), IO modules, SSRs/contactors, E-stop → \$600

LIBS mini head (OEM pulsed laser + gated spectrometer, 200–900nm) → \$800 (entry level; upgrade later)

OES spectrometer (300–900nm, fiber, lens) → \$250

RGA (bench quadrupole, 1–100amu, used/reconditioned) → \$250

## 2.6 Power & Energy (~\$1,500)

120V line conditioner/UPS or portable power station (~1.8kW cont., ~1.5kWh) ~ \$1,000  
PDU, breakers, cabling, GFCI, ground kit ~ \$300  
Optional 400-600W PV input cabling for recharge ~ \$200

Subtotal (new parts): ~\$10,800

Targeted v0.1 under \$10k strategies: source used/auction items for RGA/pump/UPS (~\$800 to ~\$1,500); choose resistance elements over induction (~\$300); use a single pyrometer (~\$150); DIY frame (~\$200). Practical v0.1 BOM ~ \$8.8-9.8k.

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## 3) Step by Step Build

### 3.1 Frame & Insulation

Build a rigid frame (80/20 extrusion or angle steel). Interior clear volume for hot zone: ~200mm x 250mm H.

Line with 1700°C fiber boards; add blanket layers; install reflective shields (Mo or W foil) with standoffs.

Add service ports: feeder entry, viewport, pyrometer peeks, electrode feedthroughs, gas ports. Keep leak paths short; use ceramic bushings.

### 3.2 Heater & Hot Zone

Option A ~ MoSi elements: Install 3-4 elements around the liner; wire to SSRs/transformer per vendor specs; keep sightlines to melt.

Option B ~ Induction: Wind water-cooled copper coil around liner zone; mount driver; ensure RF shielding/grounding.

Install primary ZrO<sub>2</sub> liner (melter) nested in a structural setter. Place W cathode tip ~10-20mm above floor sump; mount anode opposite wall (prefer inert anode or protected graphite with minimal exposure).

### 3.3 Phase Separation Geometry

Machine/fit a weir that lets light slag overflow to a slag pocket while dense Fe collects in a deep sump under cathode.

Add a Si collection pocket isolated from Fe sump by a sill. Later, a tilting ladle or tap hole can draw Si to a transfer ladle.

### 3.4 Vacuum Refining Pot

Assemble small vacuum can with internal crucible support and heater.

Fit cold trap on exhaust path; add baffles cooled by water loop. Route to pump via isolation valve; include pressure gauges at can & foreline.

### 3.5 Gas/Vacuum Skid

Mount rotary vane pump on vibration pads; plumb KF manifold to MOE cell and refining pot with isolations and check valves.

Build Argon loop: buffer tank ~ dryer ~ filter ~ mass flow controller ~ process. Return through filter to buffer.

O<sub>2</sub> vent line from MOE headspace to an exterior safe vent (or O<sub>2</sub> bottle if you choose to

capture).

### 3.6 Controls & Sensing

Install PLC, IO, SSRs/contactors; wire interlocks (door, over temp, vacuum OK).  
Mount pyrometers (two color) with purged sight tubes toward melt and refining pot.  
Mount LIBS windowed port over feeder/melt; align fiber to gated spectrometer.  
Mount OES fiber to watch electrode/plasma zone.  
Plumb RGA to refining pot foreline (through throttled tee after cold trap).

### 3.7 Power

Route mains (120V) → portable power station/UPS → PDU → heaters/controls.  
Separate clean supplies for controls; bond grounds; GFCI where appropriate.  
Verify breaker sizing (20A recommended), wire gauge, and earth bonding.

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## 4) Commissioning & First Run

### 4.1 Dry Runs

Leak check with argon; verify <50 sccm leak at 50 mbar over ambient.  
Heat soak hot zone to 1000°C; check element balance, pyrometers, thermal uniformity.  
Pump down refining pot; test cold trap condensation with water/ice first.

### 4.2 Feed Characterization

Sieve feed <250 µm; bake at 150–200°C to dry salts/moisture.  
LIBS scan (190–900 nm) over a representative sample to estimate oxide fractions (Fe oxides, SiO<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, CaO, TiO<sub>2</sub>, alkalis). Save the spectrum for the run file.

### 4.3 MOE Operation (Fe–Si–Al)

Melt/hold: Bring hot zone to 1600–1650°C under light argon (50–150 mbar over ambient).  
Fe stage: Apply low cathodic overpotential/current density; monitor OES for Fe lines. Once Fe pool accumulates in sump, tap/skim Fe or park it (mechanically isolated).  
Si stage: Step potential upward; watch Si emission grow, and OES oxygen lines. Maintain controlled headspace; ensure Fe sump temperature slightly lower to limit mixing.  
Manage bubbles; avoid vigorous stirring. If LIBS/OES signals show reduction (alloying), step back, let Fe settle, then proceed.

### 4.4 Transfer to Refining Pot

Ladle/tap molten Si into covered transfer crucible; minimize air exposure.  
Move to refining pot; close, evacuate to <10 Pa; heat to 1600–1850°C for 30–90 min.  
RGA watch: P<sub>2</sub> species decay; SiO (m/z ~44) rise indicates over evaporation → increase trapping/cool baffles or adjust temperature.

### 4.5 Casting

Pour into Si–N coated quartz mold; directional cool (1–5 K/min) from bottom to top to segregate residual impurities to the hot end; crop tail.  
Optional: mini float zone pass on a small rod for semiconductor experiments.

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#### 5) Control Logic (simplified)

Preheat to 1600°C & stable for 20min.

LIBS feed OK? If not, abort.

MOE: set Fe window & Fe sump mass threshold & tap or isolate.

MOE: set Si window & Si rate threshold met & end Si stage.

Transfer & vacuum refine & RGA P below setpoint & cool.

Cast & cool & log batch & spectra.

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#### 6) Safety

Thermal: >1700°C potential. Use shields, IR PPE, lockouts. No combustible clutter.

Gas: O<sub>2</sub> evolution; ensure venting away from personnel. No HF used; off-gas is primarily O<sub>2</sub>/Ar/SiO<sub>2</sub>; cold-trap captures condensables.

Electrical: 120V at high current; bonded grounds; GFCI; E-stop. UPS/inverter must be rated for continuous draw.

Optical: Bright hot surfaces; avoid direct viewing; use proper IR/UV eye protection when inspecting.

Mechanical: Interlocks on doors/panels; guarded auger; hot-work signage.

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#### 7) Maintenance

Inspect liners & electrodes every ~10 hours at temp; replace worn anodes.

Bake argon dryer, change filters quarterly.

Clean cold-traps each run; reclaim condensed SiO<sub>2</sub> where feasible.

Calibrate pyrometers quarterly; verify LIBS/OES wavelength daily warm-up.

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#### 8) Upgrade Path

Inert anode (ceramic-metal) to fully eliminate carbon contact.

Larger MOE pot w/ dual-zone heaters for higher throughput.

Better spectrometers for trace B/P detection (deep-UV arms).

Automated tapping, robotic ladle, and closed-loop recipe tuning from spectra.

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#### 9) Appendices

##### A) Target Setpoints (starting)

MOE hot-zone: 1625°C

Fe stage: low overpotential; current density ~0.2-0.4 A/cm<sup>2</sup> (tune by OES)

Si stage: step +10-20% vs Fe stage; watch OES (Si) & headspace

Vacuum refine: 1700-1800°C, <10 Pa, 30-90min

## B) Minimal Tools

IR thermometer, torque drivers, crimp kit, TIG level gloves, face shield, tongs, scale, sieve set.

## C) Expected Consumables (per 10 runs)

Argon: 1 lb (with reclaim)

Liners: 1 ZrO<sub>2</sub> primary + 1 Al<sub>2</sub>O<sub>3</sub> backup

Anode face: 1 (if consumable)

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v0.1 total (with used/auction gear): \$8.8k - 9.8k

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