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Technologies Carbon Capture and Storage (CCS)

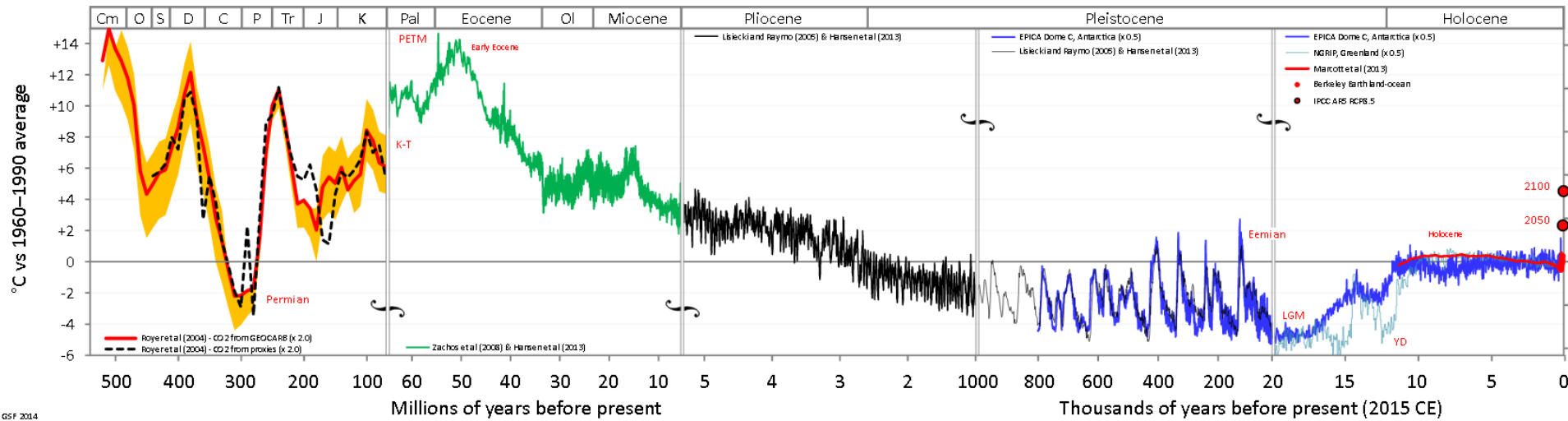
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WHY CAPTURE AND STORAGE OF CO₂?

- there is an opinion that humankind-produced CO₂ (and increase of its atmospheric concentration) is a cause of so-called global climate change or global warming
- major CO₂ sources: fossil fuel combustion, industry (steel, cement), crude oil refining
- 1860-2010 atmospheric concentration has risen from 300 to 400 ppm
- CCS currently does not belong into priorities of energy and climatic policy
- Actual State Energy Conception expects a role for CCS after 2040 and recommends a support for CCS research

Temperature of Planet Earth



CCS technologies in general

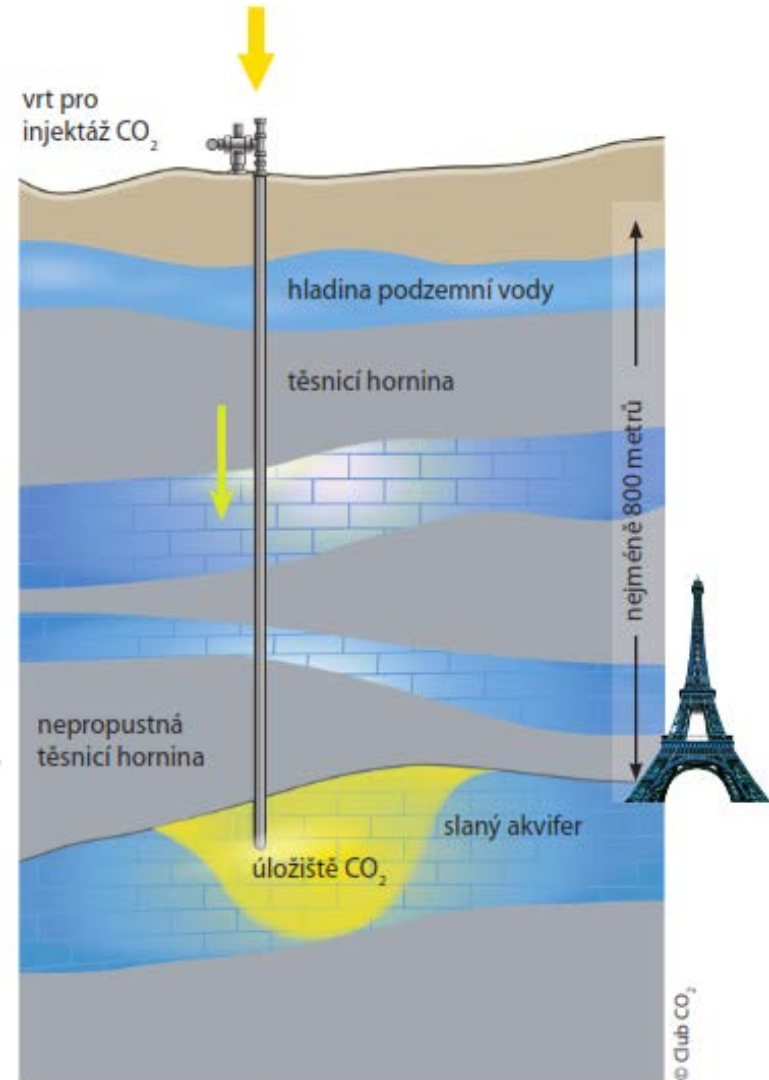
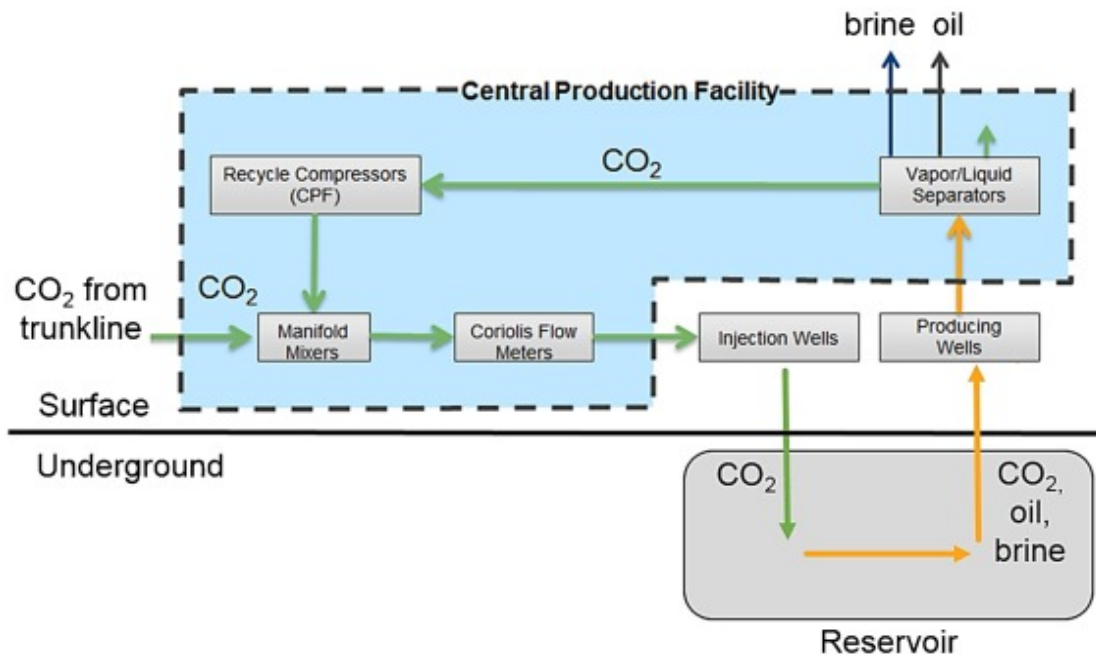
- **3 TECHNOLOGIC PARTS:** capture, transport, storage (alternatively CCU – utilization)
- **CAPTURE:** 3 main technology groups – post-combustion, pre-combustion and oxyfuel combustion
- **TRANSPORT:** gaseous or liquid, different pressures and temperatures according to economic effectiveness; longer distances – pipeline transport in gas phase, ship transport in liquid phase; shorter distances – railway transport in liquid phase
- **STORAGE:** saline aquifers – porous rocks containing saline water; exploited oil and gas fields – EOR; alternative options – deep sea hydrates, some specific rocks (e.g. basalt)

CCS technologies in general

➤ **STORAGE:** CO₂ in time (100-1000 years) reacts with surrounding environment to form new minerals

➤ **UTILIZATION:**

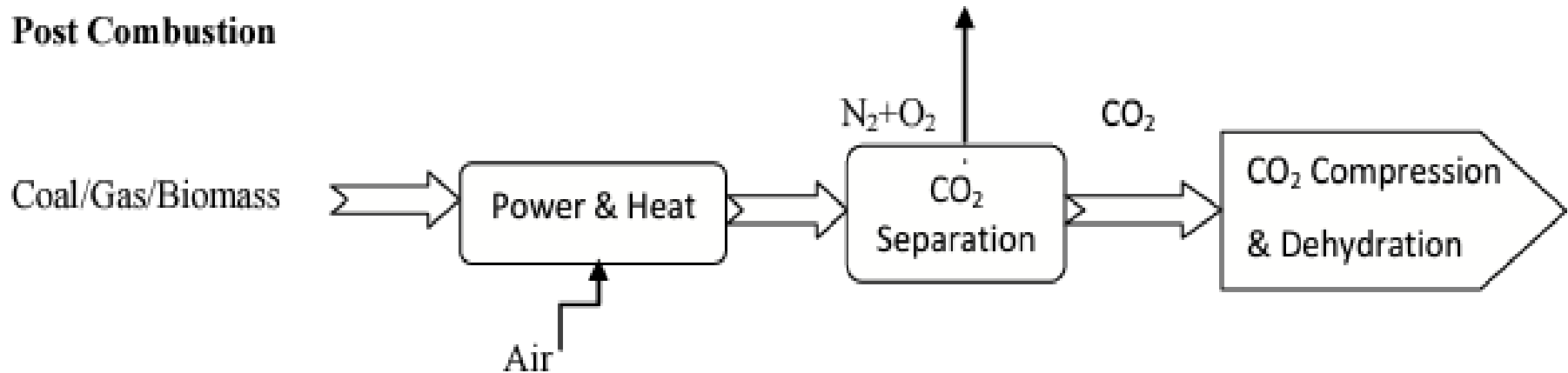
- EOR – enhanced oil recovery (and simultaneous storage)



POST-COMBUSTION CAPTURE

- = separation of CO_2 after a combustion process has been completed = capture of CO_2 in a low concentration from large gas flows
- 250 MWe unit (NW Bohemian lignite coal) produces approx. 770 thousand Nm^3/h of flue gas with CO_2 concentration approx. 14 %

Post Combustion

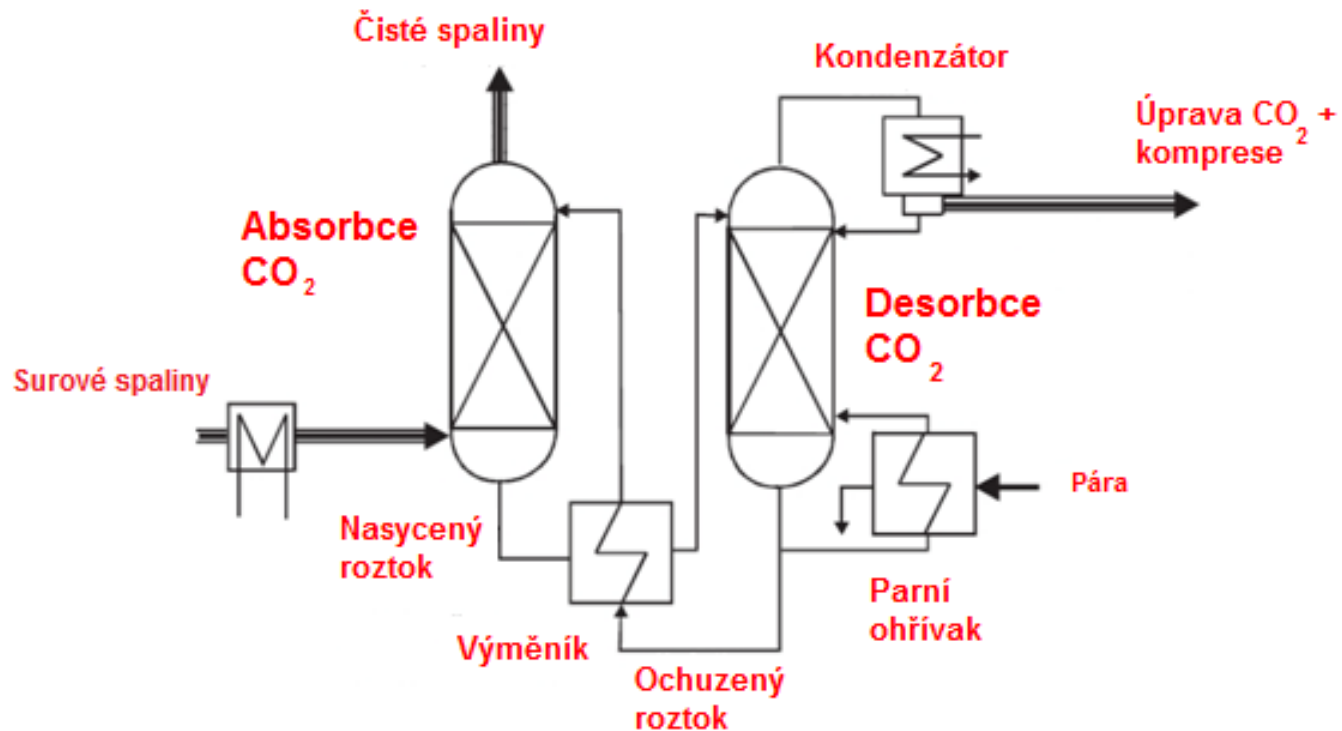
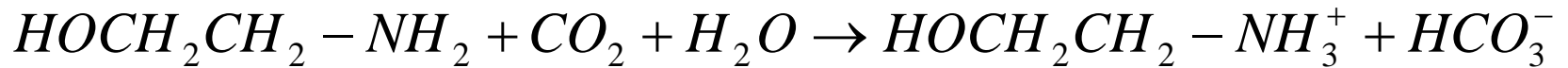


POST-COMBUSTION separation processes

Technology	Principle
Absorption	<ol style="list-style-type: none">1) Amine scrubbing, e.g. MEA or DEA2) NH_3 – ammonia scrubbing3) Alkaline solvents, e.g. NaOH
Adsorption	<ol style="list-style-type: none">1) Ca-based sorbents, e.g. limestone – Ca-looping2) Alkaline carbonates, e.g. Na_2CO_33) Special sorbents and molecular sieves, e.g. zeolites or active carbon
Membrane separation	<ol style="list-style-type: none">1) Polymeric membranes2) Inorganic and hybrid membranes
Cryogenic separation	Separation of CO_2 by freezing

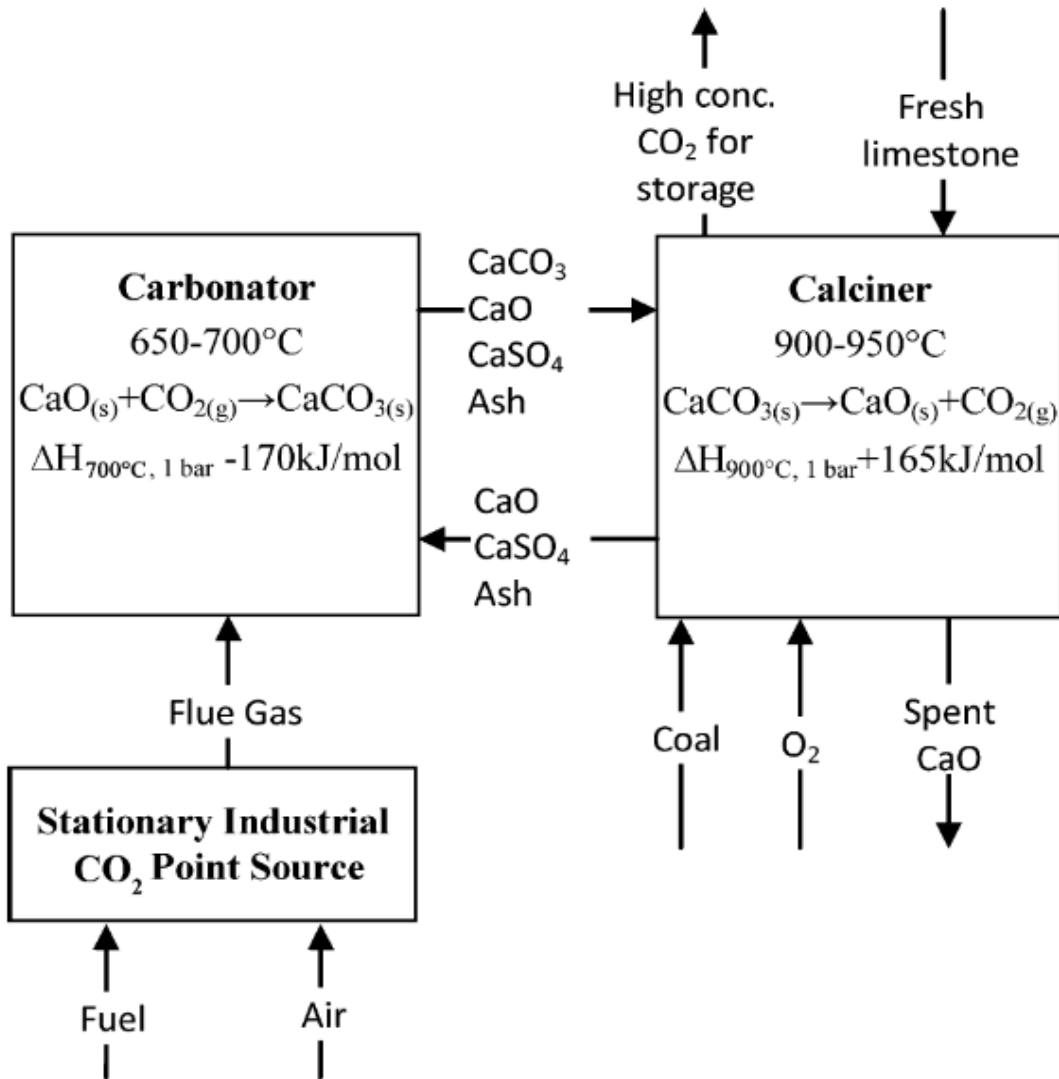
MEA SCRUBBING

- demo unit SaskPower – Boundary Dam power plant, Canada
- flue gas after deNO_x and deSO_x, regeneration in desorber; compared to ammonia: (+) higher absorption temperature (15°C); (-) lower capacity, solvent degradation



Ca-looping

➤ dual CFB principle



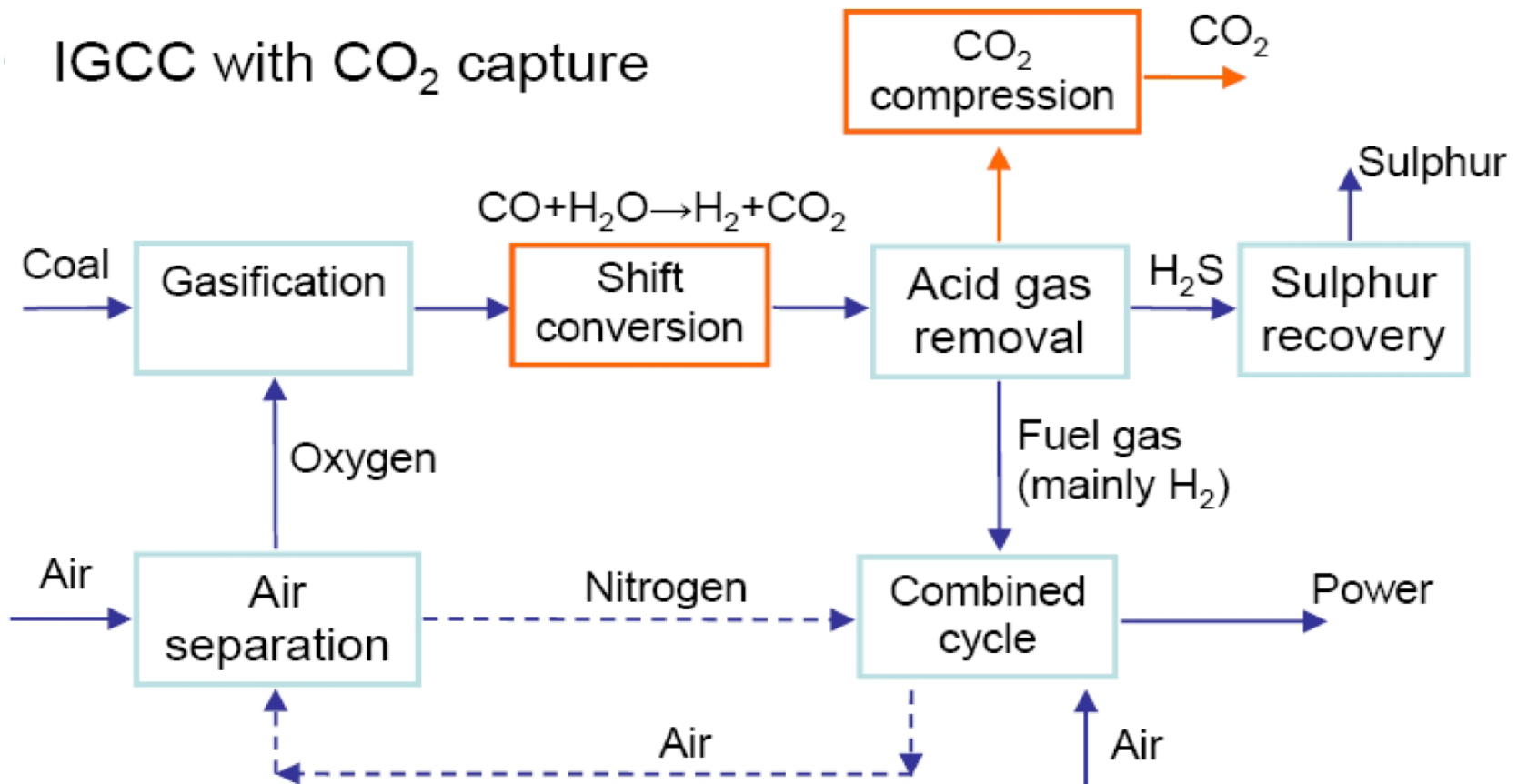
➤ Ca-looping application in a conventional power plant means **building a new source of about half thermal input in oxyfuel mode**

➤ (+) proven CFB technology, cheap sorbent, recoverable energy;

➤ (-) oxygen consumption, plant size, lower capture ratio, electricity need for driving the circulation of solids

PRE-COMBUSTION capture

- based on fuel gasification, conversion of CO to CO₂ and subsequent separation, product is H₂
- hydrogen is used in conventional IGCC



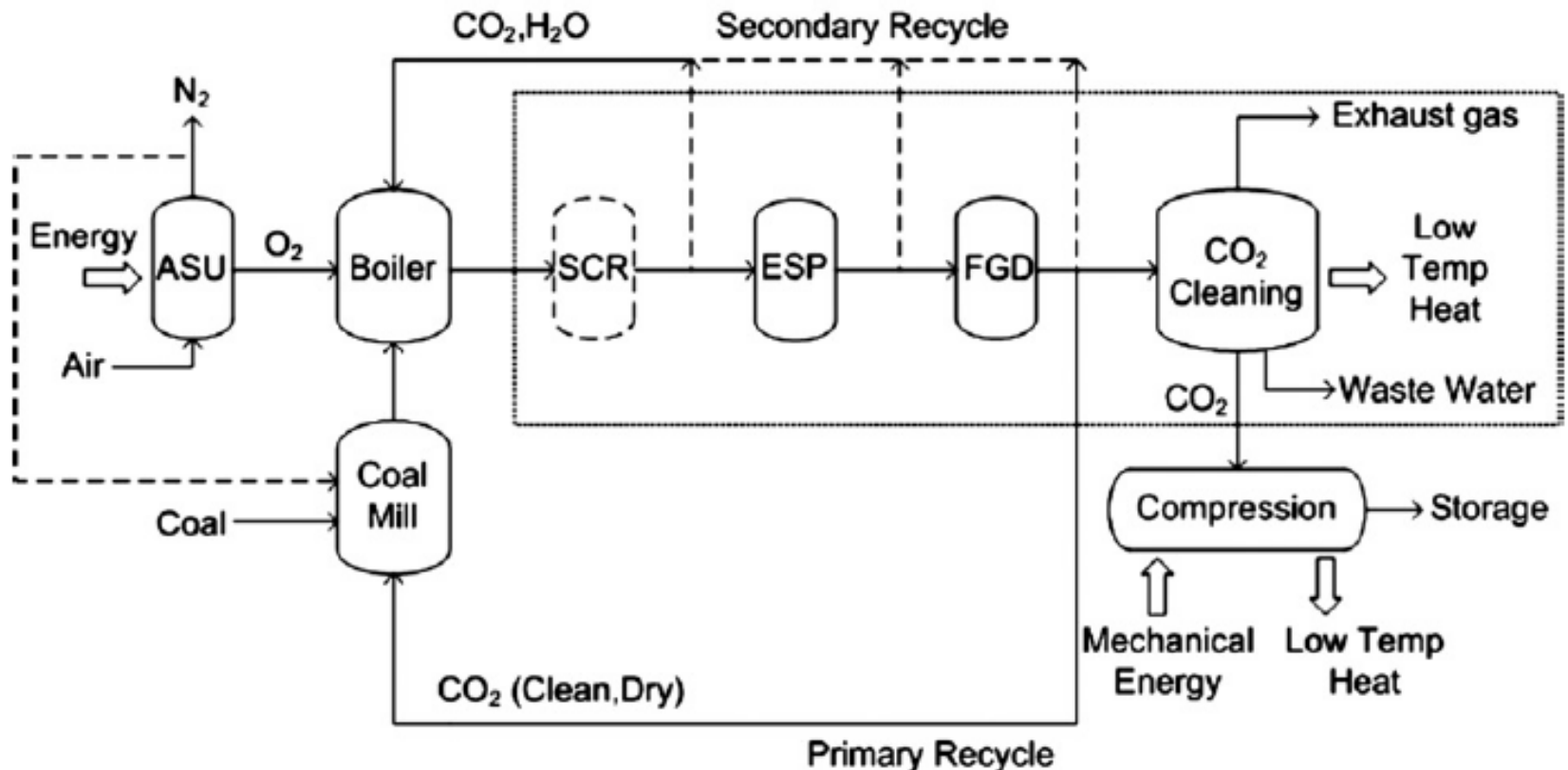
PRE-COMBUSTION capture

- Water-gas shift: two-step conversion of CO to CO₂; catalytic process (ZnO/Al₂O₃), LTS 150-300°C, HTS 350-600°C
- CO₂ is separated using an AGR (acid gas removal) technology – either with acid gases (WGS before separation) or stand-alone (WGS after separation)

Procedure	Process name	Specification
Absorption (no chemical reaction)	Selexol/Coastal AGR	Solvent mixture of dimethyl ethers and propylene glycol (DEPG); temperature 20°C
	Rectisol/Ifpexol	Solvent methanol, sub-cooled -20 to -60°C (IGCC Vřesová)
	Purisol	Solvent N-methyl-pyrrolidone (NMP), chilled to -15°C
	FluorSolvent	Solvent propylen-carbonate; chilled to -18°C
Absorption with chemical reaction	MEA	Solvent mono-ethanolamine

OXYFUEL COMBUSTION

- technologically most simple capture – combustion of a fuel with pure oxygen, output is concentrated $\text{CO}_2 + \text{H}_2\text{O}$ vapor (+minor pollutants)
- (+) simple technology, requires „only“ ASU construction; (-) purity of output CO_2 , concentration lowered by false air intake and oxygen excess; corrosion and deposits; different dimensioning of heat exchanging surfaces



COMPARISON OF CCS TECHNOLOGIES – 250 MWe reference, 9.5 MJ/kg, 90 % capture

Parameter		Current plant	NH ₃ scrub	Oxyfuel	Solid adsorbent/ zeolite	Ca-looping *)	Pre-combustion/ Rectisol*)
El. out brutto	MWe	250	238	262 **)	245	354	272
Fuel consumption	t/h	214	214	217	214	369	99**)
Internal consumption	MWe	24	75	94	47	123	41
CO ₂ production	t/h	211	211	216	211	211+113	211
CO ₂ capture	t/h	0	190	190	158	297	201
CO ₂ to air	t/h	211	21	26	53	27	10
El. out netto	MWe	226	163	168	198	231	231
Electric efficiency	%	38.4	28.12	28.53	33.73	23.01	39.75
Efficiency decrease	p.b.	0	10.87	10.46	4.77	15.39	-----

*) coal SD, calorific value 16,5 MJ/kg **) fuel pre-dried to 11% - WTA drier



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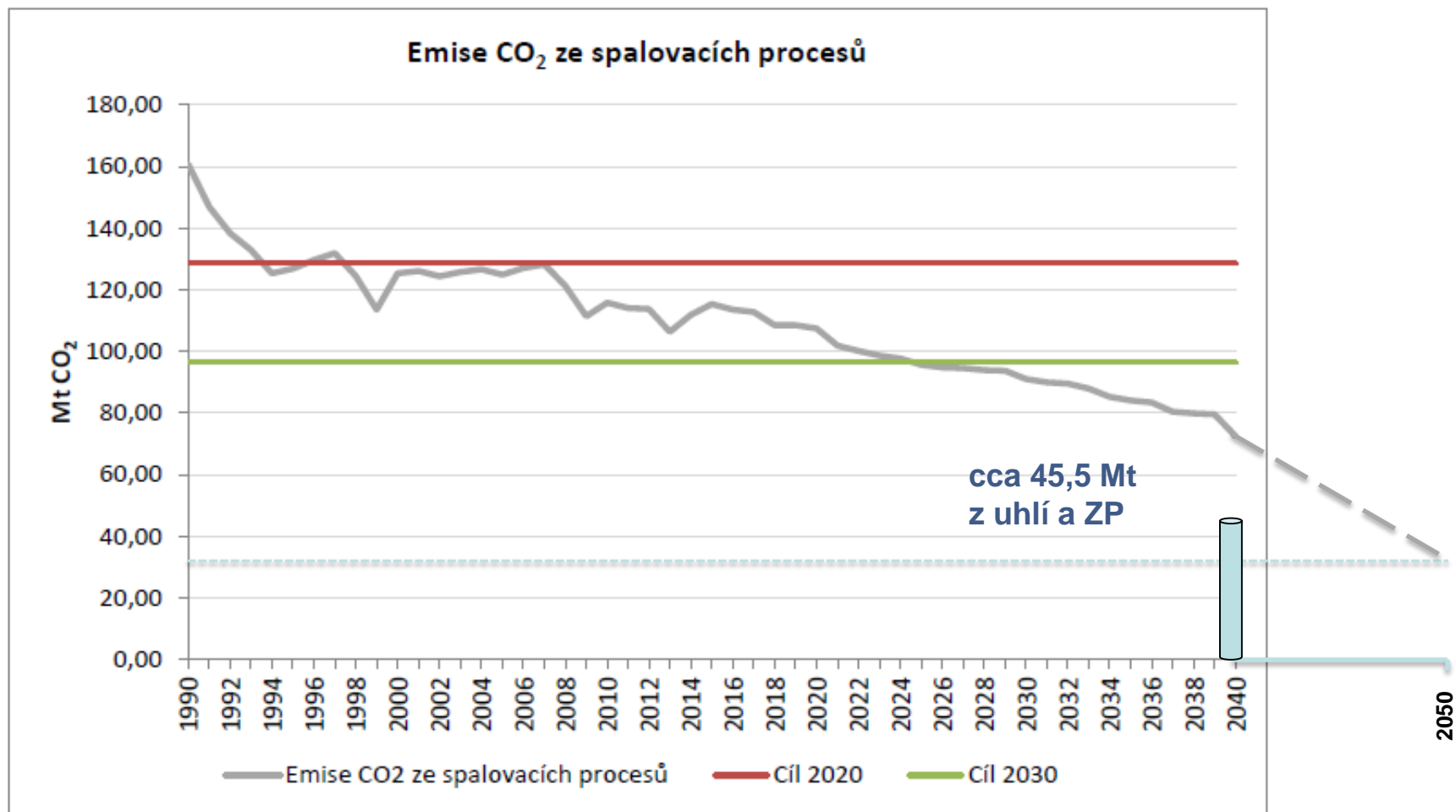
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Thank you for your attention!



CCS in the Czech Republic

Graf č. 33: Emise CO₂ ze spalovacích procesů



Pozn.: Pro emise CO₂ nejsou stanoveny cíle pro jednotlivé země EU, ale pouze cíl pro EU jako celek. Uvedené linie jsou tedy vypočteny z hodnot cíle EU snížení emisí do roku 2020 o 20 % oproti roku 1990 a cíle EU snížení emisí do roku 2030 o 40 % oproti roku 1990 vztahených k hodnotě emisí ze spalovacích procesů na území České republiky.