



CO₂ Capture by Moving Bed Temperature Swing Adsorption

Kent S. Knaebel
Adsorption Research, Inc.
6175-D Shamrock Court
Dublin, OH 43016-1200
Tel.: 614-798-9090
Fax: 614-798-9091
e-mail: k.knaebel@adsorption.com

September 27, 2008

EXECUTIVE SUMMARY

An “end-of-pipe” process has been developed to capture carbon dioxide (CO₂) from flue gas from coal-burning power plants, cement plants, and the like. It will treat flue gas immediately after the fly ash is removed. The presence of minor amounts of fly ash, however, should not pose a serious problem. The process employs temperature swing adsorption (TSA). It uses solid adsorbent to take-up CO₂ from flue gas. And it uses the heat contained in the flue gas to drive-off the enriched CO₂ from the adsorbent. The keys to performance are: the flue gas temperature and the CO₂ concentration, which are equivalent to *the available heat* and *the required heat*, respectively. Together, they dictate whether the heat contained in the flue gas is sufficient to drive the separation. When they are properly balanced, the need for parasitic energy is minimal. In contrast, the conventional alternative means for CO₂ capture is scrubbing, which requires about 30% of the gross output of a power plant’s steam in order to regenerate the scrubbing solution. Hence, the need for parasitic energy is enormous when scrubbing is used.

The ARI TSA CO₂ capture process has been evaluated via experiments and computer simulations. They revealed important parameters, e.g. the amount of adsorbent required and the size of the equipment, which represent the majority of the capital cost. The ARI TSA process tests showed that both high CO₂ purity (99%) and high recovery (89%) were possible. Additional tests with CO₂, H₂O, and SO₂ in a simulated flue gas showed no signs of adsorbent breakdown. A patent application has been filed (20060230930), and a PCT application (Publication No.=WO 2006/112977) was submitted.

Adsorbent attrition is of minimal concern on account of features that allow less breakage and friction than for other moving bed designs, and a device by which adsorbent fines (if they occur) can be continuously removed.

The design is modular, i.e., the system will be comprised of dozens of identical parallel units. Each one could be amenable to shut-down for maintenance, without adversely affecting the processing capacity of the remainder. If sufficient redundancy is provided for, plant operation and output should not be impacted.

Now that preliminary testing and engineering are complete, we plan to build a pilot-scale TSA process (1 ton of CO₂ per day) to evaluate the process under realistic conditions. The project is being organized by ADA-ES, based in Littleton, Colorado. These tests will yield engineering data and will assess its technical and economic feasibility. The cost for testing will be partly covered by a DOE grant that was recently announced, along with industrial contributions, and will require about 2 years.

PROCESS DESCRIPTION

The process is illustrated below. This diagram was taken from an animated version. As such, it represents a specific instant of its operation, during which the adsorbent, gas, bucket conveyor, and a heat-transfer medium are all moving.

Let's start with the adsorbent beads, being delivered to the top of the adsorption section by the bucket conveyor. At this point the beads (blue) are cool, and relatively free of CO₂. As the beads pass over a screen, any fines that may have been generated (white dots) during adsorption or regeneration are separated. The whole beads fall onto a series of perforated trays. The perforated trays are designed to retard the beads, so that their residence time in the adsorption section is a few minutes. While in the adsorption section, the beads contact flue gas, consisting of adsorbable components such as CO₂ (shown as red squiggles) and non-adsorbable components such as nitrogen (shown as yellow squiggles). Flue gas enters the adsorption section through the perforated distributor pipe at the bottom of that section. The adsorbable components are taken-up the adsorbent as they pass countercurrently, and the non-adsorbable components escape through the stack at the top.

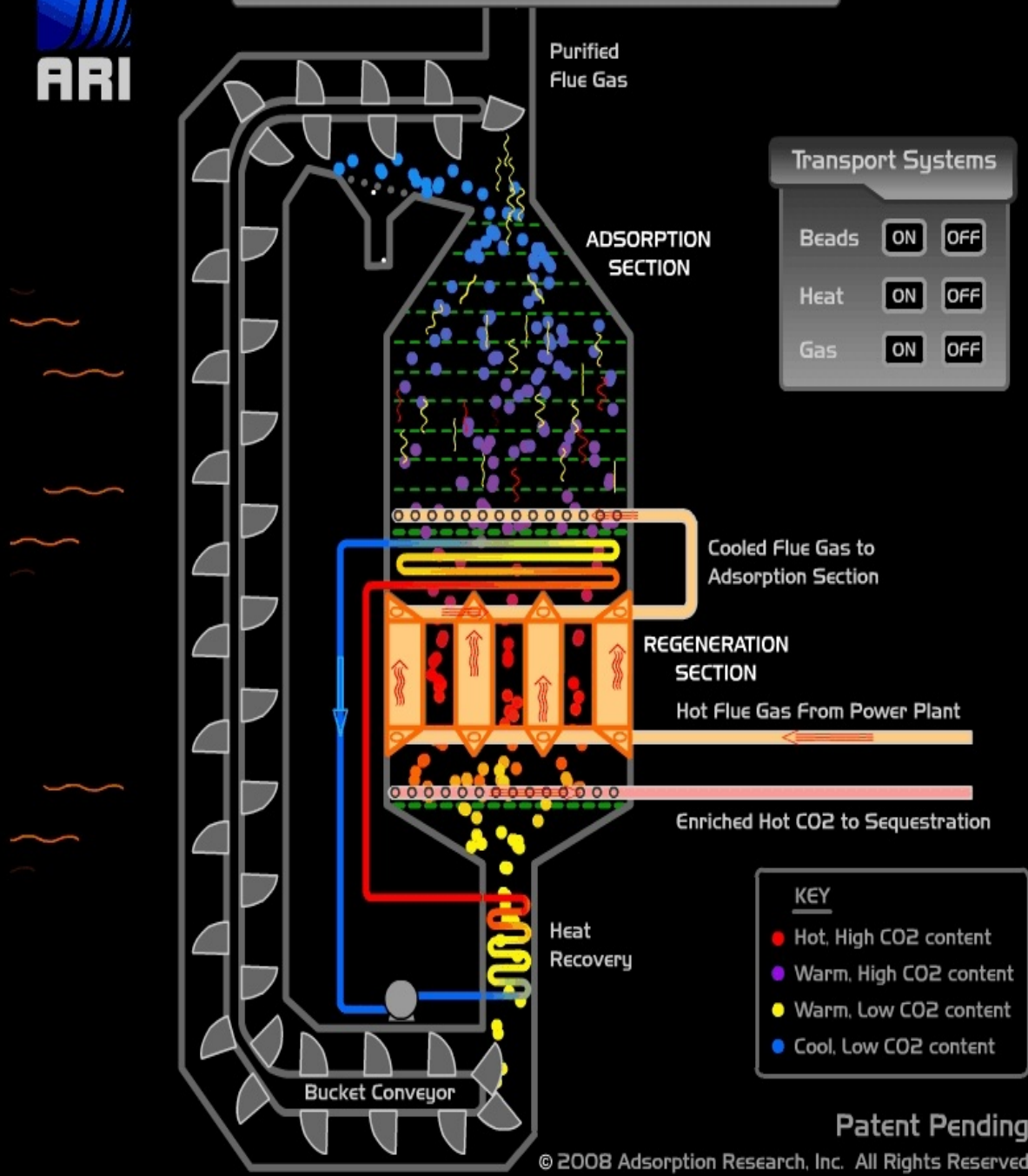
Upon reaching the bottom of the adsorption section, the adsorbent beads are fully loaded, and they proceed to the regeneration section. First, the beads are preheated, as they pass through a countercurrent heat exchanger. In that section, the beads pick-up heat, which is recycled from beads in the heat-recovery section. (The latter beads are hot and relatively free of CO₂.) The beads then contact a heat exchanger section in which the hot flue gas delivers its heat to the beads. Now the beads are red. As can be seen, the hot flue gas enters this heat exchanger section from the power plant, in a duct (at the right-hand side). The flue gas and adsorbent bead remain separated by the heat exchange surface, which is just a metal barrier. The diagram shows the flue gas in a rectangular duct, with triangular sections at the bottom and top, respectively, through which the hot flue gas is delivered and the relatively cool flue gas collected. (The relatively cool flue gas may subsequently be quenched in a separate vessel – not shown.) That relatively cool flue gas is then introduced to the adsorption section through the perforated distributor pipe mentioned earlier. The hot beads readily desorb the CO₂ and any other adsorbable components, and their color changes to yellow. Meanwhile, the enriched CO₂ is collected and withdrawn in a perforated pipe, to sequestration.

As mentioned earlier, the hot adsorbent beads, depleted of CO₂ are cooled by the circulating fluid in the heat recovery section. Afterwards, the cooled beads fall into buckets of the conveyor and are lifted back to the top of the adsorption section. As they are lifted, additional cooling occurs, indicated by the brown horizontal squiggles, which emanate from the bucket conveyor enclosure, at the left.

A Key identifies the states of the adsorbent beads according to colors at various locations. The block labeled "Transport Systems" refers to animation features, which obviously are all "on" in the snapshot. Those allow one to view the various aspects individually, or in pairs, and to appreciate the relative motion and synchronization, which of course is not possible with a still image.



ARI's Moving-Bed TSA CO2 Capture Process



Justification

The laboratory data and computer simulations were used to generate approximate capital and operating cost estimates of a commercial CO₂ recovery system. In addition, budgetary quotes were obtained from vendors to arrive at an overall plant cost estimate. The main design parameters and results are shown in Table 1.

Table 1. Operating and Capital Costs for CO₂ Capture by the ARI TSA Process at a 500 MW Coal Fired Power Plant.

Flue Gas Flow Rate:	=	1,500,000 SCFM = 2,200 MMSCFD
CO ₂ Emission Rate	=	15,000 tons per day
Flue Gas CO ₂ Concentration	=	12% (vol) from the plant
Flue Gas H ₂ O Concentration	=	12% (vol) from the plant
Outlet Gas CO ₂ Concentration	=	1.20% (vol) from the TSA system
Min. CO ₂ Recovery Percentage	=	92%
Max. Exhaust Gas Pressure Drop	=	<5 in. H ₂ O in the TSA system
Adsorbent Reservoir	=	10,000,000 lb
Temperatures		
Flue Gas Inlet Temp.	=	500°F
Outlet Gas Temp.	=	125°F
Adsorbent Regeneration Temp.	=	350°F
Dimensions		
Approximate Footprint and Height	=	300 ft x 200 ft x 50 ft.
Estimated Cost		
Fabricated Steel	=	\$147 million
Initial Adsorbent Inventory	=	\$ 10 million
Building & Foundation	=	\$ 3 million
<u>Installation + Contingency</u>	=	<u>\$262 million</u>
Approximate Capital Cost	=	\$420 million.
Amortization (6% x 15 yr)	=	\$ 43 million per year
Power Cost	=	\$ 7 million per year
Total Processing Cost*	=	\$9.65 per ton of CO ₂

* Assuming the system is on-line 95% of the time, i.e., 347 days per year.

As can be seen, the estimated capital cost includes the structural steel and adsorbent, as well as the building, foundation, installation labor, and contingency. The resulting operating cost per ton of CO₂ captured will depend on the accuracy of these estimates, as well as the amortization rate and term. This is the cost for the captured CO₂, at 1 atm, without drying (which we expect to cost less than an extra \$1 per ton).