TSC Jitterbug Fusion of D in Pd NanoClusters
Frank Dodd (Tony) Smith Jr. - 2015

viXra 1501.0234

Abstract:
Graphic Cover summarizes TSC Fusion of D in 147-atom Palladium Clusters; Jitterbug Icosa - Cubocta - Icosa Transformation reloading D into Pd Clusters; and Zeolite Steam Engine utilizing Fusion Energy.

The body of the paper gives details of the Pd - D TSC Jitterbug Fusion Process:

Clusters of Palladium atoms (also clusters of atoms of Nickel and similar elements) have two basic structures:
Icosahedral and Cuboctahedral

1 - Icosahedron <-> Cuboctahedron Jitterbug Transformation

2 - Pd clusters with absorbed Deuterium have two states:
   Icosahedral ground state
   Cuboctahedral metastable state

3 - Tetrahedral Symmetric Condensation (TSC) in Icosahedral Pd-D produces Fusion.

4 - Icosahedra TSC Fusion Triggers Jitterbug to Cuboctahedra.

5 - Cuboctahedra reload TSC sites and Jitterbug back to Icosahedra

6 - Repeat the Cycle:
Table of Contents:

Graphic Cover ... page 1

1 - Introduction ... page 4

2 - Geometry of TSC Fusion within Palladium Cluster loaded with Deuterium ... page 9

3 - Root Vectors and the Jitterbug Transform ... page 14

4 - Structure of Palladium Clusters ... page 15

5 - Deuterium Loading of Palladium Clusters and Jitterbug Cycle time ... page 19

6 - Jitterbug Transformation between Icosahedron and Cuboctahedron ... page 20

7 - Jitterbug D-Loading analogy with Browning M1911 and Maxim Gun ... page 23

8 - TSC Fusion of 4D to 8Be* to 2 alpha giving 47.6 MeV ... page 24

9 - TSC 47.6 MeV energy release mechanisms ... 27

10 - Parchamazad experiments: D-loaded Palladium in Sodium Zeolite Y ... page 28

11 - a TSC-Jitterbug Fusion device with 1 milligram of Palladium in the form of 147-atom Pd clusters could produce 100 KiloWatt-Hours in an hour ... page 33

12 - Sandia 1.5 nm Pd Clusters ... page 34

13 - ITQ-37 Zeolite with pore 2 nm ... page 37

14- TSC-Jitterbug-Zeolite Fusion energy ... page 38

15 - TSC-Jitterbug-Zeolite Fusion energy as Heat and Steam ... page 40

16 - TSC-Jitterbug-Zeolite Fusion energy as Ultracapacitor charge ... page 43

17 - Global Energy and TSC-Jitterbug-Zeolite Fusion machines ... page 45

NOTE - Most of the detailed quantitative discussion in this paper is for Pd / D Fusion but Nickel / Hydrogen is very similar to Palladium / Deuterium so the same processes may hold for Pd / H and Ni / D and Ni / H Fusion.
**Introduction:**

D = Deuterium = Heavy Hydrogen nucleus has 1 Neutron + 1 Proton

D + D has 2 Neutrons + 2 Protons

He = Helium nucleus also has 2 Neutrons + 2 Protons

BUT

( using Einstein’s E = mc² )

**D+D is heavier than He by 23.8 MeV**

( where 1 MeV of energy = about 2 x mass of electron )

so

**Fusion of D+D to He produces Energy**

**How much Energy ?**

Earth’s Oceans contain a lot of Water, most of which is H₂O = HHO

where H = Hydrogen nucleus = 1 Proton

but

0.015% = about 1 part per 7,000 of Earth’s Ocean Water is D₂O = DDO

D₂O can be extracted by processes such as distillation and chemical exchange.

If 1 / 1,000 of the Earth Ocean D₂O were used for Fusion of D+D to He

the Energy would be enough to sustain

10 billion people consuming energy at the level of 1985 USA

for 2,000,000 years

**How much would it Cost ?**

The energy equivalent of a gallon of D₂O is about equal to 300,000 gallons of fuel oil.

You can buy a liter of D₂O for about $1,000

which is about 1 cent per gallon of oil (energy equivalent).
Can D+D to He Fusion be done without using the High Energy Input needed by Hot Fusion?

Around 1999 at Osaka University, Arata and Zhang filled a tube of Palladium metal with Palladium-black which is Palladium particles of size about 5 nanometers and injected high-pressure Deuterium gas producing. for some batches of Palladium-black, Energy and He continuing as long as pressure was maintained.

Why did only some batches produce Energy? My guess is that the 5-nanometer average size of Palladium particles is not the optimal size for D+D to He Fusion Energy but some batches may contain a lot of the optimal size which I think is about 1.5 nanometers.

What is so special about 1.5 nanometers? 1.5 nanometers is the size of a cluster of 147 Palladium atoms which has two states, an Cuboctahedral Metastable State and an Icosahedral Ground State, that can transform into each other by a Buckminster Fuller Jitterbug Transformation that facilitates the Fusion Energy Process.
Has anyone gotten consistently reproducible results?

Around 2012 at University of LaVerne, Iraj Parchamazad experimented with Deuterium gas injected into a chamber containing Palladium embedded in Zeolites producing heat **Energy in 10 out of 10** runs of the experiment with Energy produced on the scale of **kiloWatts of power per milligram of Palladium**.

Each Zeolite Cage contains a Palladium atom cluster. 

**In the Iraj Parchamazad experiments,** the Zeolite cage size is 1.3 to 2.4 nanometers and each Palladium Cluster contains on the order of the optimal 147 atom = 1.5 nanometer size.
Even though experiments show that D+D to He Fusion works with Palladium Clusters, Hot Fusion needs High Temperature / Energy for two Deuterium nuclei both of which are have Positive Electric Charge = Coulomb Barrier repelling each other so

**How can Deuterium nuclei in Palladium Clusters get close enough for Fusion?**

Julian Schwinger said:

“... in the very low energy cold fusion, one deals essentially with a single state, described by a single wave function, all parts of which are coherent ...”.

Akito Takahashi uses a **single coherent state of 4 Deuterium nuclei and 4 electrons** within Palladium Cluster structure for a **Tetrahedral Symmetric Condensate** process:

The 4 D = D+D+D+D and 4 electrons are each in Tetrahedral Symmetry. Their single coherent state has net Electric Charge = 0, so there is no Coulomb Barrier to prevent them from Condensing together close enough for the D+D+D+D to Fuse producing an excited Beryllium Be nucleus with 4 protons and 4 neutrons which then splits into two He nuclei plus 2 x 23.8 = 47.6 MeV Energy, much of which Energy is carried away by the 4 electrons when they interact with Palladium Cluster electrons and the Zeolite electric field.
How can Jitterbug Icosa-Cubocta Transformations of 147-atom Palladium Clusters facilitate Tetrahedral Symmetric Condensate D+D+D+D to Be to He+He + Energy Fusion?

147-atom Palladium Cluster with absorbed Deuterium is in Icosa Ground State and TSC Fusion heats the Palladium cluster and the Zeolite.

The heated Palladium cluster expands by Jitterbug to the Cubocta Metastable State and the Fused Deuterium nuclei are replaced by new Deuterium from the gas.

The Cubocta Metastable state, loaded with the new Deuterium, collapses by Jitterbug back to the Icosa Ground State.

Then a new TSC Fusion occurs, and the cycle repeats as long as there is Deuterium “ammunition” to fuel the Fusion.
Geometry of TSC Jitterbug Fusion of Deuterium in Palladium Clusters

Akito Takahashi has developed a Tetrahedral Symmetric Condensate (TSC) model for fusion D+D+D+D -> 8Be -> 4He + 4He + 47.6 MeV in Pd atomic clusters. The same type of process may occur with Ni instead of Pd and/or H instead of D.

This paper describes the geometry of Pd atomic clusters and how it enables TSC fusion of D within the Pd clusters.

A 147-atom Palladium cluster has two special states:

- A metastable cuboctahedral state
- An icosahedral ground state

The icosahedral 147-atom ground state has 12 exterior vertices and a central icosahedron with 12 interior vertices which are the innermost and outermost vertices of 12 exterior TSC Fusion sites.

There are 13 TSC Fusion sites (12 of which are exterior) in each 147-atom Pd Cluster.
This image (adapted from a Thingiverse web site image) shows the 12 Palladium nuclei (blue dots) arranged in an icosahedral configuration plus a 13th Palladium nucleus at the center of the configuration that is the TSC fusion site (orange dot fusion site containing blue dot Palladium nucleus). Within the icosahedral configuration the 4 Deuterium nuclei (red dots) and their 4 electrons (green dots) are at the vertices of a cube and are in a single coherent quantum state.
That state then contracts in a symmetric way along the cyan lines going down to the center of the cube at the central Palladium nucleus (blue dot) which acts as a quantum attractor, combining into the Deuterium TSC quantum state to guide the condensation of the quantum state of Deuterium nuclei and electrons.

The D + D + D + D does Fusion at the orange dot to form 8Be which decays to 4He + 4He + 47.6 MeV energy most of which energy is carried away by the 4He+4He and the 4e electrons of the TSC coherent quantum state.
The 4e electrons may carry, by quantum processes, most of the energy to the Palladium cluster electrons and on to the Zeolite in which the Palladium cluster is enclosed.

Some of the energy goes to a Jitterbug transformation of the icosahedral Palladium, depleted of Deuterium fusion fuel, to a cuboctahedral configuration which has 6 large square openings through which ambient Deuterium Fuel can enter the Palladium cluster.
After entering the Palladium cluster the 4 Deuterium nuclei (red dots) and 4 electrons (green dots) form a Tetrahedral Symmetric Coherent Quantum State centered on the 8 triangular faces of the cuboctahedral configuration. Then, since the icosahedral configuration is the Palladium cluster ground state, another Jitterbug transformation takes the Palladium cluster to an icosahedral configuration with the replenished Deuterium nuclei and electrons ready for another round of TSC fusion.
Root Vectors and the Jitterbug Transform

12 vertices of a Cuboctahedron (image from Fulton and Harris) are D3 = A3 Root Vectors with Weyl Group of order 4x6 = 4x3x2 = 24

The Cuboctahedron transforms by Jitterbug into an Icosahedron (image from Wikipedia)

Icosahedron = Root Vectors of H3 (Humphreys)
“... Crystallographic groups ... stabilize... a Lattice ...
H3 and H4 ...[are]... non-crystallographic groups ...
H3 is the symmetry group of the Icosahedron ...”.
H3 has Weyl Group of order 120 = 5x24 = 5x4x3x2.
Structure of Palladium Clusters

There are two basic structures that are Jitterbug Transforms of each other:

**Icosahedral and Cuboctahedral**

- **n** = number of shells
- **N** = number of Pd atom vertices
- **d** = diameter of icosahedral configuration in nm
- **C** = number of cells in icosahedral phase
- **CT** = number of tetrahedral cells in icosahedral phase
- **CO** = number of octahedral cells in icosahedral phase

<table>
<thead>
<tr>
<th>n</th>
<th>N</th>
<th>d</th>
<th>C = CT + CO</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>1</td>
<td>0.27</td>
<td>0 = 0 + 0</td>
</tr>
<tr>
<td>1</td>
<td>13</td>
<td>0.70</td>
<td>20 = 20 + 0</td>
</tr>
<tr>
<td>2</td>
<td>55</td>
<td>1.13</td>
<td>100 = 80 + 20</td>
</tr>
<tr>
<td>3</td>
<td>147</td>
<td>1.56</td>
<td>280 = 200 + 80</td>
</tr>
<tr>
<td>4</td>
<td>309</td>
<td>2.00</td>
<td>(icosa and cubo images not shown)</td>
</tr>
</tbody>
</table>

How many TSC Fusion sites are in a 147-atom Pd cluster?

The icosahedral 147-atom ground state has 12 exterior vertices and a central icosahedron with 12 interior vertices which are the innermost and outermost vertices of 12 exterior TSC Fusion sites.

(see viXra 1502.0069)

There are 13 TSC Fusion sites (12 of which are exterior) in each 147-atom Pd Cluster.

**How do the Icosahedral Clusters grow to 147 atoms?**


"... The Mackay icosahedron is obtained by packing tetrahedra and octahedra around an icosahedron [12 vertices +1 center = 13 vertices]...

if an octahedron is placed on every face of an icosahedron, the angular gap between neighboring octahedra can be closed by a very small deformation, to bring them into face contact [12 + 20 x (6-3)/2 = 42 vertices]...

... The concave regions of the resulting polyhedron can
be filled by five-rings of tetrahedra \([42 + 12 + 1 \text{ center} = 55 \text{ vertices}]\)...
What about more than 147 atoms?

As more layers are added, the deformations of tetrahedra and octahedra accumulate and eventually destabilize the structures necessary for Jitterbug and TSC Fusion. The next Mackay cluster beyond 147 atoms has $147+162 = 309$ atoms.

Barretau, Desjonqueres, and Spanjaard in Eur. Phys. J. D. 11 (2000) 395-402 say: “... the icosahedron is the preferred structure at small sizes, and the critical size at which the relative stability becomes favorable to cuboctahedrons is $N = 561$ for PdN clusters [for which]...
For $N = 13$ the cuboctahedron is ... unstable.

For $N = 55, 147$, and 309 atoms the cuboctahedron is metastable and slightly distorted. Its transformation to a perfect icosahedral structure needs an activation energy of 12 meV for $N = 55, 28$ meV for $N = 147$ and 45 meV for $N = 309$. The activation energies involved in the inverse transformation are 61 meV for $N = 55, 51$ meV for $N = 147$ and 48 meV for $N = 309$.

...[ compare 47.6 MeV for each TSC Fusion event ]...

$N = 309$ is disfavored for TSC-Jitterbug Fusion with respect to $N = 147$ for two reasons:

energy levels are too close for rapid Jitterbug cubocta to icosa transition

$N = 309$ Pd Cluster is too large (2 nm) to fit through 1.5 nm expanded Sodium Zeolite Y pore
It seems that **147 atoms is optimal for TSC fusion.**

**How many D/H atoms can live in a 147-atom Pd/Ni cluster?**

F. Calvo and A. Carre say in Nanotechnology 17 (2006) 1292–1299 "Structural transitions and stabilization of palladium nanoparticles upon hydrogenation": "... Cuboctahedra ...[and]... icosahedra ... contain exactly the same number of atoms. ... In the case of ... the 147-atom Pd cluster ... the favoured structure in the pure metal is the three-layer icosahedron.

![Figure 1. Palladium clusters fully loaded with hydrogen. (a) Pd\textsubscript{147}H\textsubscript{200}, I\textsubscript{h} symmetry; (b) Pd\textsubscript{147}H\textsubscript{164}, O\textsubscript{h} symmetry.](image)

Since the minimum full load for Icosa or Cubocta Pd/Ni 147-atom clusters is 164 D/H atoms, no more than 3 cycles of full TSC fusion (each consuming 56 D/H nuclei) can occur without replenishment of D/H from the surroundings of the clusters (such as immersion of the clusters in D/H gas).

**How long does it take Deuterium to load into Palladium?**

Wang, Hara, and Watanabe in Materials Transactions, Vol. 48, No. 3 (2007) pp. 560 to 565 say: "... Pure Pd, Pd-4 at%Pt and Pd-8 at%Pt ... powders smaller than 200 mesh (<74 mm) were prepared ... hydrogen absorption ...[by Pd-4 at%Pt]... was extremely fast and attained to equilibrium within tens of seconds. Hydrogen absorption by Pd and Pd-8 at%Pt was also very fast ...".

**Tens of seconds is much longer than the times for TSC Fusion and for Jitterbug** so it determines the time duration of one TSC-Jitterbug Fusion Cycle and for the purpose of rough calculations it seems reasonable to take

$$36 \text{ seconds} = 1/100 \text{ hour} = \text{time duration of one TSC-Jitterbug Fusion Cycle.}$$

This time is much shorter than the usual loading time for old-type Cold Fusion experiments using Palladium rods, discs, much-larger-than 1.5 nm powder, etc because there are only 3 layers of Pd atoms in 1.5 nm 147-atom Pd clusters.
Jitterbug Transformation between Icosahedron and Cuboctahedron

Icosahedra and Cuboctahedra both have 12 vertices so that it is possible to transform them into each other. Buckminster Fuller called that transformation the Jitterbug.

To make Cuboctahedra (unit edge length) from Icosahedra (unit edge length) choose 6 pairs of Icosahedra triangle faces (white in the above images) and lengthen the common edge of each pair by a factor of \(\sqrt{2}\). That expansion flattens each of the triangle pairs to produce 6 square faces of the Cuboctahedron. The other Icosahedral \(20 - 2\times6 = 8\) (shaded) triangle faces are rotated and become the other \(14 - 6 = 8\) triangle faces of the Cuboctahedron.

, thus increasing the number of faces from \(8 + 6 = 14\) to \(8 + (6 + 6) = 20\) while keeping the number of vertices constant at 12.

There are two ways to choose a diagonal of an Icosahedron triangle face pair in the construction, corresponding to the two possible orientations of an Icosahedron.

Choice of diagonal for one Icosahedra triangle face pair forces (by requiring consistency) the choices for all other face pairs of all Icosahedra.
The triangle faces of the Icosahedron/Cuboctahedron are rotated by a Golden Ratio angle defined by sliding Icosahedron vertices on the edges of a circumscribing Octahedron from points dividing edges into Golden Ratio segments to points dividing edges into two equal segments so that the Octahedron then circumscribes a Cuboctahedron. If the edge lengths of the Icosahedron/Cuboctahedron are kept the same then the Octahedron surrounding the Cuboctahedron will be an expansion of the Octahedron surrounding the Icosahedron.

Just as in the choice of a Cuboctahedron square diagonal to be compressed, there are two ways in which the edge could be divided into Golden Ratio segments, corresponding to the two possible orientations of an Icosahedron. Choice of Golden Ratio segments for one edge forces (by requiring consistency) the choices for all other edges.


The volume expansion of the Jitterbug Transformation from Icosahedron (unit edge) to Cuboctahedron (unit edge) is:

Icosahedron volume = \((5/12) \times (3 + \sqrt{5})\) = 2.18169499
Cuboctahedron volume = \((5/3) \times \sqrt{2}\) = 2.3570226

Icosahedron/Cuboctahedron volume ratio = \(0.9256147947\)
Cuboctahedron/Icosahedron volume ratio = \(1.0803630254\)
How do Jitterbug Transformations enable D reloading?

The cuboctahedral configuration resulting from Fusion Energy Jitterbug is not only larger than the Icosahedral configuration, it has 6 large square openings allowing easier entry into the Pd cluster of the Deuterium Nuclei (red dots) and Electrons (green dots).

Also, the cuboctahedral configuration has 8 small triangle faces to which the 4 Nuclei and 4 Electrons of the Deuterium are attracted to form the Tetrahedral Symmetric Coherent Quantum State which is maintained after the second Jitterbug to the icosahedral ground state thus producing a reloaded icosahedral configuration for another round of TSC fusion.
Jitterbug D-Loading analogy with Browning M1911 and Maxim Gun

(with Colt Series 80 Government 10 mm Delta Elite version of Browning's M1911 semi-auto)

"... The M1911 ... use[s] ... the short recoil ... action ... Cycle ...
1. Ready to fire position. [Slide] is locked to barrel, both are fully forward.
   [Icosahedral Pd with D atoms in TSC positions]
2. Upon firing, [slide] and barrel recoil backwards a short distance while locked together. Near
   the end of the barrel travel, the [slide] and barrel unlock.
   [Firing = D-D Fusion]
3. The barrel stops, but the unlocked [slide] continues to move to the rear, ejecting the empty
   shell and compressing the recoil spring.
   [Recoil Spring = Icosahedral Stability Phase induces transformation of Cuboctahedra]
4. The [slide] returns forward under spring force, loading a new round into the barrel.
   [Loading New Round = Cuboctahedral D atoms moved to Icosahedral TSC positions]
5. [Slide] locks into barrel, and forces barrel to return to battery.

... The very first short-recoil-operated firearm was also the first machine gun, the Maxim gun.
... Vladimirov also used the short recoil principle in the Soviet KPV-14.5 heavy machine gun. ..."
(quote from Wikipedia entries on M1911 pistol and on Recoil operation)
TSC Fusion of 4D to 8Be* to 2 alpha giving 47.6 MeV


Every particle in TSC can make central squeezing motion with same velocity, to keep charge neutrality of total TSC system ... this squeezing motion can be treated as Newtonian mechanics until when four deuterons get into the range (about 5 fm) of strong nuclear interaction. ... TSC starts Newtonian squeezing motion to decrease linearly its size from about 100 pm radius size to ... the minimum size state ... as shown in ... Semi-classical view of squeezing motion of TSC, \(<e> = (e_↓ + e_↑)/2\) for QM view at four electron centers ...

[Note that the TSC process is spontaneous not requiring initial stimulus.]
... Classical squeezing motion ends when four deuterons get into the strong force range (5 fm) and/or when four electrons get to the Pauli's limit (about 5.6 fm for e-e distance). Here for Pauli's limit, we used the classical electron radius of 2.8 fm. Since the range of strong interaction is comparable to the classical electron diameter (5.6 fm) ... the intermediate nuclear state $^8\text{Be}^*$ will be formed just after the minimum size state ...

Immediately at ... $^8\text{Be}^*$ formation ... 4d-cluster shrinks to much smaller size (about 2.4 fm radius) of $^8\text{Be}^*$ nucleus, and four electrons should go outside due to the Pauli's repulsion for fermions. Shortly in about few fs or less (note; Lifetime of $^8\text{Be}$ at ground state is 0.67 fs), $^8\text{Be}^*$ will break up into two $^4\text{He}$ particles ...[ with energy released of about 47.6 MeV, mostly as photons of a few keV energy each ] ...

... when TSC is just formed ... averaged electron position (electron center of \( <e> = \frac{\text{e}_\downarrow + \text{e}_\uparrow}{2} \), Bosonized electron pair ...) ... locates at vertexes of regular cube with tetrahedral combining orbits and outer dilute clouds ...

... At ... cube ... vertexes ... three Bohr wave functions superpose and electron density is about nine times larger than that of outer dilute cloud. Therefore, the semi-classical treatment of central squeezing motion by Newtonian is approximately fulfilled for "coherent"central averaged momentums for eight particles. ...

As soon as 4D/TSC(t=0) state with D2 molecule size (Rdd = 74 pm) is formed ...
the QM-Langevin equation gives numerical solution for time-dependent $R_{dd}$ and mean relative kinetic energy of d-d pair of a face of 6 TSC (d-e-e-d-type) faces, as copied from reference and shown in Fig.10. ...

... The ‘adiabatic’ size of 4D/TSC reaches at a few tens fm size in 1.4 fs, so fast. With adiabatic 4D/TSC size around 20 fm, 4D-fusion takes place by ...

$$D + D + D + D \rightarrow 8\text{Be}^*\ (E_x = 47.6\text{ MeV}; J^*)$$ ...

Fusion yield per 4D/TSC generation is calculated by integrating time-dependent fusion rate by the Fermi’s first golden rule ... that was very close to 1.0, namely 100%, during the very small time interval of ca. $2 \times 10^{-20}$ s in the final stage of condensation.

Mean relative kinetic energy of neighboring d-d pair of 4D/TSC-minimum is ca. 14 keV, which is accidental resembling value to the hot fusion experimental devices as ITER (DT plasma).

... the quantitative study on the TSC formation probability in D(H)-loaded metal systems is yet to be done by solving many-body time-dependent problems under organization field of condensed matter. It is challenging work ...".
TSC 47.6 MeV energy release mechanisms

As Takahashi said “... Immediately at ... 8Be* formation ...
4d-cluster shrinks to much smaller size (about 2.4 fm radius) of 8Be* nucleus, and four electrons should go outside due to the Pauli’s repulsion for fermions. Shortly in about few fs or less (note; Lifetime of 8Be at ground state is 0.67 fs), 8Be* will break up into two 4He particles ...

There are at least 3 possibilities for energy release from 8Be* (Ex = 47.6 MeV):

1 - Schwinger Coherent Electron process:
Julian Schwinger in 1990 lecture at Universite de Bourgogne said: “... in the very low energy cold fusion, one deals essentially with a single state, described by a single wave function, all parts of which are coherent ...”. That is the basic idea of Takahashi’s TSC process.
At the time of 8Be* formation by TSC, instead of the “... four electrons ... go[ing] outside ...” the four electrons should remain part of Schwinger’s “coherent ... single state” and some of the 47.6 MeV should go to the four electrons.
If the 47.6 MeV were to be shared equally by the two 4He and the four electrons, then each 4He alpha particle and each electron would carry about 8 MeV. After decoherence of the TSC Fusion state, the 4He alpha particles would remain as measurable Fusion Products and the electrons would join the Pd metal electrons, thus heating the Pd Cluster.

If the Pd Clusters were embedded in Zeolite Cages, heat from the Pd Cluster would be transferred to the Zeolite, from which it could be extracted by the Zeolite-Water process.

Further experiments are needed to show whether or not this is what really happens.

2 - Each of the two 4He alpha particles carry away about 23.8 MeV.
Experiments by Roussetski indicate that this does not occur.

3 - Takahashi’s Nucleon-Halo BOLEP (burst of low energy photons) process producing 1 - 10 keV soft X-rays.
Akito Takahashi said in a September 2014 email message: “... my recent theory of nucleon-halo model (JCF13, attached) maximum alpha-particle energy from 8Be* by 4D-fusion is 17 MeV … 23.8 MeV alpha particles should not be emitted either by the 4D-fusion or by the DD fusion ...”.
Takahashi’s paper JCF13 says
“... The $^8\text{Be}^*$ (Ex = 47.6 MeV) may damp its excited energy by major BOLEP (burst of low energy photons) process from $<n-h-h-n>$ nucleon-helion halo state ...

... to $^8\text{Be}$-ground state ... A complex decay scheme is proposed ...

Major decay channel is modeled as an electro-magnetic transition of BOLEP to the $^8\text{Be}$-ground state which breaks up into two 46 keV alpha-particles ...

BOLEP is modeled as emission of ... stochastic burst events of ca. 1.5 keV averaged energy photons ...

Minor channels are modeled as BOLEP transitions to lower ... states (Ex = 34, 27.5, 22.98, 22.0, 20.1, 16.6, 11.4 and 3.04 MeV), from where two-alpha break-up channels open ... emit[ting] ... alpha-particles at 17, 13.8, 11.5, 11, 10, 8.3, 6.9, 5.7 and 1.55 MeV ... which meets ... with observed data by Roussetski et al ...

The asymmetric break-up from the Ex = 34 MeV state has a branch to emit 5.2 MeV triton, which will induce secondary D-t reaction ... to emit 9-19 MeV (En) neutrons ...

Further experiments are needed to show whether or not this is what really happens.
Parchamazad experiments: D-loaded Palladium in Sodium Zeolite Y

The size required for Jitterbug / TSC Fusion is a Palladium atomic cluster whose ground state is icosahedral and can easily Jitterbug Transform into a cuboctahedral state and whose size is large enough to contain several TSC Fusion Cluster sites, each of which is an icosahedron that can Jitterbug transform into a cuboctahedron.

The 13-atom Pd/Ni cluster (0.70 nm) is an icosahedron, for 1 TSC Fusion Cluster site.

The 2-shell 55-atom Pd/Ni cluster (1.13 nm) has two icosahedra that share a central vertex, for only TSC Fusion Cluster sites.

Clusters of between 56 and 147 atoms contain from 2 to 13 TSC Fusion Cluster sites by partially filing the 3rd shell of atoms.

The 3-shell 147-atom Pd/Ni cluster (1.56 nm) has 12 exterior TSC Fusion Cluster sites plus 1 central TSC Fusion Cluster sites, so it contains 13 TSC Fusion Cluster sites.

Clusters of between 147 and 309 atoms contain at least 13 TSC Fusion Cluster sites.

The 4-shell 309-atom Pd/Ni cluster is 2.00 nm in size, so it is disfavored with respect to the 3-shell 147-atom cluster for use with Sodium Zeolite Y whose pore size is 0.74 nm expandable to 1.5 nm.

Iraj Parchamazad has experimented with Palladium embedded in Zeolites, getting reproducible heat production in amounts consistent with TSC-Jitterbug Pd-D Fusion:

$$47.6 \text{ MeV} \times 14 \text{ TSC Sites} / 147\text{-atom Pd Cluster} \times 4.45 \times 10^{-17} \text{ Watt-Hours / MeV} = 2.965 \times 10^{-14} \text{ Watt-Hours / 147-atom Pd Cluster for each Jitterbug Cycle}$$

Mass of 147-atom Pd Cluster 147 \times 106 \times 1.66 \times 10^{-21} = 2.587 \times 10^{-17} \text{ milligrams}

so a milligram of 147-atom Pd Clusters gives about 1 KiloWatt-Hour each Cycle.
Iraj Parchamazad uses **Sodium Zeolite Y** also known as faujasite. The Wikipedia page for faujasite says:

“... The faujasite framework consists of sodalite cages which are connected through hexagonal prisms. The pores are arranged perpendicular to each other. The pore, which is formed by a 12-membered ring, has a relatively large diameter of 7.4 Å [0.74 nm] The inner cavity has a diameter of 12 Å [1.2 nm] and is surrounded by 10 sodalite cages. ...”.

Ruby Carat and Melvin Miles interviewed Iraj Parchamazad of University of La Verne in 2012. In that video interview Iraj Parchamazad said that the Zeolite cavity size can oscillate and vary, enlarging up to about 2.4 nm.

A corresponding enlargement of pore size is to about 1.5 nm which would permit a 3-shell 147-atom Palladium cluster to enter the Zeolite Cavity.

Iraj Parchamazad did not use Sandia’s 1.5 nm Palladium clusters in his Zeolite but used an organometallic solution containing Palladium atoms. After putting that into the Zeolite he heated the Zeolite to burn off Carbon, Hydrogen, and Oxide leaving a Zeolite and some Palladium. Then he exposed the Zeolite/Palladium to Deuterium, and got excess heat 10 out of 10 times, indicating fusion.
As to how much Palladium was put into the Zeolite, he found

To use those results to see how many Palladium atoms were in each Zeolite Cavity, look at the detailed structure of Sodium Zeolite Y.

Each Zeolite Cavity is surrounded by 10 sodalite cages which are arranged in a 3-dimensional Diamond network as shown in this image

from
The Zeolite Cavity has geometric symmetry related to an isometric trapezoid with the 4 holes corresponding to deleting 4 octahedral-type 3-face groups.

The 10 red dots correspond to the 10 sodalite cages.

Each sodalite cage is a 24-vertex truncated octahedron as in this image from Wikipedia.

A more detailed view of a sodalite cage from http://som.web.cmu.edu/structures/S099-sodalite.html has red dots for Oxygen and pink spheres for OH and yellow spheres for Sodium and blue tetrahedra for Silicon and green tetrahedra for Aluminum.

It shows that of the 24 vertices of the sodalite cage, 12 are Aluminum and 12 are Silicon so each Zeolite Cavity has $10 \times 12 = 120$ Aluminum atoms.
To see the number of Aluminum atoms in each Zeolite Cavity look at Iraj Parchamazad’s graph of the weights of Aluminum and Palladium and consider that their Atomic Weights are 27 for Aluminum and 106 for Palladium. Looking at the graph, it appears that the weights are about 7 for Aluminum and 32 for Palladium so the number of Palladium atoms in each Zeolite Cavity is about

\[ N_{pd/z} = 120 \times \left( \frac{32}{7} \right) \times \left( \frac{27}{106} \right) = 140 \text{ atoms} \]

which is roughly equal to the number of atoms (147) in a 3-shell Pd atomic cluster

The 3-shell 147-atom icosahedral Palladium atomic nanocluster contains 14 TSC Fusion Sites and each TSC Fusion event produces 47.6 MeV

\[ 47.6 \text{ MeV} \times 14 \text{ TSC Sites} / 147\text{-atom Pd Cluster} \times 4.45 \times 10^{-17} \text{ Watt-Hours / MeV} = 2.965 \times 10^{-14} \text{ Watt-Hours / 147-atom Pd Cluster for each Jitterbug Cycle} \]

Mass of 147-atom Pd Cluster 147 \times 106 \times 1.66 \times 10^{-21} = 2.587 \times 10^{-17} \text{ milligrams}

so that a milligram of 147-atom Pd Clusters can produce roughly 1 KiloWatt-Hour in each TSC-Jitterbug Fusion Cycle

If 36 seconds = 1/100 hour is taken as the Cycle time then

a TSC-Jitterbug Fusion device with 1 milligram of Palladium in the form of 147-atom Pd clusters with D-Loading should produce 100 KiloWatt-Hours in an hour.
147-atom Pd clusters have diameter about 1.5 nanometers.
1.5 nm Pd Clusters have been produced
at Sandia National Laboratories
and University of New Mexico Center for Micro-Engineered Materials
according to a Journal of Catalysis article
"Facile, surfactant-free synthesis of Pd nanoparticles for heterogeneous catalysts"
at
by Patrick D. Burton, Timothy J. Boyle, and Abhaya K. Datye showing

Tim Boyle said in email October 2014:
“... We easily remade the Pd NP just need to get TEM to see what size they are.
If they come out good, we can go ahead and make some for you.
Couple of things.
This is very easy and ya'll may want to do it yourselves
(esp after the next couple of comments).
Simply dissolve Pd-acetate in MeOH and stir for 5 min,
let grow for 20 more and should have your size.
The problem is these will continue to grow and plate out onto the sides of the container,
unless you use a substrate.
Would you want these on a substrate, then that'll need to be supplied.
If we make it, we'd have to send it as a solution ...
could you handle this and could you use it?
It won't be a powder, which I think is what you want.
We can dry it down to a powder but not sure what size that will be
or how they’d cluster and how they'd redisperse or in what solvent.
we can try to deposit the materials on a number of surfaces and just let it dry.
Again, not sure how the clustering of these particles will occur.
A gram will take about 2.5 g of Pd(Oac)2 which we have but will need replaced. ...”.
In April 2015 Tim Boyle of Sandia National Lab produced Palladium nanoclusters on SBA-15 mesoporous silica substrate.

In May 2015 Mark Goorsky of UCLA did XRD (x-ray diffraction), SEM and TEM (scanning and transmission electron microscope studies of the Sandia material, saying:
“... Analysis of:
tjb.Pd/SBA.30b Pd30
tjb.Pd/SBA.20b Pd20
tjb.Pd/SBA.10b Pd10
tjb.Pd/SBA.40b: the vial for this sample (40b) was broken in transit to UCLA ...
measurements were not performed on it. ...

... All samples (Pd10, 20, and 30) showed a peak near 20° 2 theta which is typically related to amorphous SiO2.
Pd20 exhibited scattered intensity near 40° 2 theta
Neither Pd10 nor Pd30 showed this scattered intensity ...
The simulation shows the broadening for the (111) and (200) Pd diffraction peaks
This broadening matches well with the presence of the scattered intensity in Pd20
The simulated structure is based on 1.2 nm particle size
1.5 nm particle size produced a peak that was slightly sharper than shown in the figure
... the SEM does not show signs of pores ...
but ... ~ 1 micron particle (SBA-15) are indeed observed ...
TEM is needed to determine presence of Pd ... nanoparticles in ... pores ...
Particles would be just below resolution of this ...[SEM]... system ...

The June 2015 TEM report for Pd30 shows some Pd but only in a few regions
which is consistent with the XRD results in which the Pd peak was not observed.

The June 2015 TEM report for Pd20 shows Pd distributed through most of the SBA-15.
The image below shows 6 sites (red circles) within a 1-micron square (green box)

4 of the red sites (1,3,5,6) were shown by EDX to be Palladium clusters.
At least 2 other black dots in the green box look like they are likely to be Palladium clusters,
so I estimate that each square micron of a layer of the material contains 5 Pd clusters.
Since there are 1,000 microns in a millimeter,
there should be 5 x 10^6 Pd clusters in a square millimeter of a layer of the material.
If the layers are each about 1 micron thick
then a cubic millimeter of the material should contain about 5 x 10^9 Pd clusters.

Since each TSC fusion event produces 47 MeV fusion energy,
how much fusion energy would be produced by 1 TSC fusion cycle in all of the Pd clusters?
Each 147-atom Pd cluster contains 12 outer icosahedra + 1 central icosahedron.
I estimate (guess) that all of the outer 12 icosahedra will produce a TSC fusion event
in each TSC fusion cycle.
There are about 4 x 10^(-17) Watt-Hours per MeV, so the total energy output for 1 TSC cycle
should be about
47 x 4 x 10^(-17) x 12 x 5 x 10^9 = 11,280 x 10^(-8) = 1 x 10^(-4) Watt-Hours = 0.1 milliWatt-Hours
For every 100 TSC fusion cycles, one cubic millimeter of the material
should produce by TSC Deuterium fusion about 10 milliWatt-Hours of energy.
I do not know the time of a TSC fusion cycle
but if 36 seconds = 1 / 100 hour is taken as the TSC fusion cycle time
then the power of one cubic millimeter of the material would be about 10 milliWatts.
I would like to see experiments with Zeolite directly using Sandia 1.5 nm Palladium NanoClusters.

If there is difficulty with getting the Sandia Clusters to fit into the Sodium Zeolite Y then I would like to see experiments with Zeolite ITQ-37 which has pore size about 2 nanometers. (Royal Society of Chemistry, 29 April 2009 and Sun et al, Nature 2009)
Pd D Zeolite Y Fusion

1 - 15 ml of methanol (MeOH) in a scintillation vial

2 - Add 5 mg palladium acetate (Pd(OAc)2) whose color is red-orange

3 - Reduce the Pd(OAc)2 by MeOH to Pd atoms by stirring for 5 minutes with unobstructed exposure to room lighting.

4 - Add 10 mg of 30-40 nm Sodium Zeolite Y Crystals in colloidal suspension. Each Zeolite Crystal will be Tetrahedral in shape. At 30-40 nm size each will have about 12 to 16 large Cavities per edge. About half of the Cavities will be on the Exterior Surface of the Tetrahedral Crystal where they will be easily accessible by Pd atom clusters in the colloidal suspension. (Microcrystalline synthetic faujasite)

   ( ? where can 30-40 nm Zeolite Y be purchased and how much does it cost ? )

5 - Place on elevated stir plate and allow to react undisturbed for 20 minutes. During 20 minutes the Pd atoms form clusters that grow to size 1.5 nm (147 atoms)

   Initially the Pd atom clusters are very small (only a few atoms) and will migrate into the large Cavities of the Zeolite Crystals and continue to grow to size 1.5 nm (147 atoms) at 20 minutes. Color of colloidal suspension changes from pale yellow to dark green over the 20 min

6 - At 20 minutes Pd-loaded Zeolite (and any remnant Pd still in colloidal suspension) are removed and the Pd-loaded Zeolite is dried

7 - Pd-loaded Zeolite is placed in reaction chamber where it is exposed to Deuterium gas from tank and calorimeter measurements are taken to measure any heat that might be produced by TSC-Jitterbug fusion. (analogous to heat produced by Arlata and Zhang (replicated by McKubre at SRI) with no external power input - only palladium powder + deuterium gas)
After Palladium TSC - Jitterbug Zeolite fusion has been established the next step is production of useful amounts of energy which energy can be used as **Heat using Zeolite-Water properties**

“... We report ... Monte Carlo simulations of water ... ... adsorption in [ Zeolite ] NaY ... faujasite ... The existence of cyclic water hexamers ... located in the 12-ring windows ... 

... recently disclosed by neutron diffraction experiments ... were ... observed in the case of NaY ...”. ( Angela Di Lella, Nicolas Desbiens, Anne Boutin, Isabelle Demachy, Philippe Ungerer, Jean-Pierre Bellat, and Alain H. Fuchs, Phys. Chem. Chem. Phys. 8 (2006) 5396-5406 )

( see viXra 1502.0096 )

or as **Electric Charge of a Zeolite-based UltraCapacitor:**
Heat

According to a 7 June 2012 techthefuture.com web article by Tessel Renzenbrink:
“... Zeolite is a mineral that can store up to four times more heat than water ... zeolite retains a hundred percent of the heat for an unlimited amount of time ... When water comes into contact with zeolite it is bound to its surface by means of a chemical reaction which generates heat. Reversely, when heat is applied the water is removed from the surface, generating large amounts of steam. The transference of heat to the material does not cause its temperature to rise. Instead, the energy is stored as a potential to adsorb water. The ... [German Fraunhofer Institute]... scientists used these particular properties to turn zeolite into a thermal storage system. They created a storage device and filled it with zeolite pellets. To charge the pellets, they exposed them to heat. To retrieve the energy they simply added water. ...”.

Here is my design for a TSC-Jitterbug Zeolite Pd-D fusion heat engine:

![Diagram of TSC-Jitterbug Zeolite Pd-D fusion heat engine](image)

(D2O Heavy Water is used to take heat from the Zeolite to make steam so that Hydrogen from H2O does not poison the TSC-Jitterbug process by replacing Deuterium in the Palladium nanoclusters, a possible problem pointed out by Melvin Miles. D20 heavy water from Fisher Scientific costs about $1,000 per liter for 99.8 atom % D.)
Preparation of Sodium Zeolite Y has unit cell size about 2.5 nanometers which corresponds to the edge-length per cavity of its overall tetrahedral structure.

According to http://www.google.com/patents/US20040047803
“... Synthesis and stabilization of nanoscale zeolite particles ...
Zeolite Y is of great interest ... Zeolite crystals prepared under conventional synthesis conditions frequently have a mean particle size of between 1 and 5 μm. ... it would ... be useful if the zeolite particles were sufficiently small to form a colloidal suspension ...
Mono- or di-saccharides can be used to keep the crystal size of faujasite (zeolite X and Y) small ...
... Sucrose, dextrose or other saccharides are added to a conventional aluminium silicate reaction mixture obtained by mixing aqueous alkali metal silicate and alkali metal aluminate solutions at low temperatures, followed by ageing and hydrothermal synthesis. Crystal sizes of between about 30 and 40 nm are claimed ...
”.

According to a Journal of Catalysis article by Patrick D. Burton, Timothy J. Boyle, and Abhaya K. Datye "Facile, surfactant-free synthesis of Pd nanoparticles for heterogeneous catalysts"
“... room temperature reduction of Pd(OAc)2 in MeOH is slow enough to produce a suspension of ... metal-phase ... Pd NPs. ...
A Pd-NP/C catalyst was prepared by mixing the carbon support into the suspension of Pd NPs and evaporating the solvent. Aggregate formation was a concern, as there were no capping agents to prevent particle growth. Therefore, the nanoparticles were collected quickly before substantial aggregation could occur. ... this technique is general and can be extended to other powder supports. ...
”.

An “other powder support” that would be useful for TSC-Jitterbug fusion energy would be 30-40 nanometer Zeolite Y Crystals in colloidal suspension.

As the Pd nanoclusters “... grow for 20 ... min ...” up to size 1.56 nm for the 147 atom size that is optimal for TSC-Jitterbug fusion, they are small enough to fit into the Exterior Cavities of the Zeolite Y Crystals ( which are have average pore opening 0.74 nm and cavity size 1.2 nm but which sizes can oscillate to be up to about twice those sizes )

Due to the open structure of the Zeolite Y Crystals, growth up to the 147 atom size can continue inside the Exterior Cavities of the Zeolite Y Crystals.

As soon as the Pd nanoclusters have grown to the 147 atom size the solvent can be evaporated and the powder of 30-40 nm Zeolite Y Crystals loaded with Palladium can be collected and placed in the TSC-Jitterbug Fusion Reaction Chamber for exposure to Deuterium gas and heating the Zeolite Y Crystals by fusion energy.
As to how many of the 30 nanometer Zeolite Y Crystal Cavities are Exterior (and therefore easily accessible to the Pd nanoclusters in colloidal suspension)
the Zeolite Y Crystal has tetrahedral structure and each unit cell with 1 Cavity is 2.5 nanometers so a 30 nm Zeolite Y Crystal would have 30 / 2.5 = 12 cavities per edge and its tetrahedral structure would have 12 triangular layers

and a total of 1+3+6+10+15+21+28+36+45+55+66+78 = 364 Cavities
of which 1+3+6+10+15+21+28+36 = 120 would be Interior Cavities (purple dots)
and 364 - 120 = 244 would be Exterior Cavities (orange dots)
so 244 / 364 = 67 % of the Cavities of the 30 nm Zeolite Y Crystals would be Exterior and therefore relatively easily accessible to the Pd nanoclusters in the colloidal suspension.

A 40 nm Zeolite Y Crystal would have 40 / 2.5 = 16 cavities per edge with 16 triangular layers
and 1+3+6+10+15+21+28+36+45+55+66+78+91+105+120+136 = 816 Cavities
of which 1+3+6+10+15+21+28+36+45+55+66+78 = 364 would be Interior Cavities
and 816 - 364 = 452 would be Exterior Cavities
so 452 / 816 = 55 % of the Cavities of the 40 nm Zeolite Y Crystals would be Exterior.

As to how much Zeolite Y Crystal should be mixed in colloidal suspension with Palladium nanoclusters that grow to 147-atom size:

The atomic mass of a Zeolite Y unit cell is
( from nptel.ac.in Introduction to Catalysis Lecture 36 Zeolites )
( 56x23 Na + 56x59 AlO2 + 136x60 SiO2 + 264x18 H2O ) = 17,504
for actual mass = 17504 x 1.66 x 10^(-21) = 2.906 x 10^(-17) milligrams

The atomic mass of a 147-atom Pd nanocluster is
147x106 = 15,582
for actual mass = 15,582 x 1.66 x 10^(-21) = 2.587 x 10^(-17) milligrams

so if all the Cavities of Zeolite Y Crystal were External
equal masses of Palladium and Zeolite Y Crystal would be optimal
but for 30-40 nm Zeolite Y Crystals only about 2/3 to 1/2 of their Cavities are External so
it may be optimal to use Zeolite Y Crystal mass = 2 x Palladium mass in the colloid.
UltraCapacitor

According to a 26 January 2011 PhysOrg.com article by Lisa Zyga:
“... The unique 3D array of nanopores in zeolite-templated carbon ...

( image modified to show
Zeolite-Templated Carbon framework as Black, Palladium NanoClusters as Purple,
Ambient Fluid with Deuterium for TSC-Jitterbug Reloading as White,
Soft X-Rays from TSC-Jitterbug Fusion as Cyan arrows
that ionize parts of the Zeolite, ejecting electrons (Blue) into the Ambient Fluid
and leaving positive ions (Red) in the Zeolite-Templated Carbon
thus building up a Capacitance Voltage
between the Zeolite-Templated Carbon and the Ambient Fluid )

... enables it to be used as an electrode for high-performance supercapacitors that have
a high capacitance and quick charge time ... The zeolite-templated carbon consists of
nanopores that are 1.2 nm in diameter ... and that have a very ordered structure ...”.

Synthesis of Zeolite-Templated Carbon is described in the 2013 Caltech Ph.D. Thesis of Nicholas Stadie
“... Zeolite-templated carbon (ZTC) materials were prepared ... by ... established
methods ...
The ZTC capacitor process converts TSC-Jitterbug fusion energy directly to electricity. Since it does not require the Zeolite-heat-water-steam chemical structure only the Zeolite Y Crystal geometric configuration is needed so all the Zeolite Y can be converted to ZTC carbon configurations attached to a single base carbon substrate that acts as a Capacitor Electrode. (It would be difficult to use separated Zeolite Y Crystals as an electrode.)

However, the ZTC has fewer Exterior Cavities than the colloidal free-floating Zeolite Y Crystals because each tetrahedral structure is attached to the carbon substrate by a base face, thus eliminating 1/4 of the Exterior Cavities so that for ZTC Capacitor fusion energy:

A 30 nm ZTC structure would have $30 / 2.5 = 12$ cavities per edge and its tetrahedral structure would have 12 triangular layers and a total of $1+3+6+10+15+21+28+36+45+55+66+78 = 364$ Cavities of which $1+3+6+10+15+21+28+36+78 = 198$ would be Interior Cavities and $364 - 198 = 166$ would be Exterior Cavities (orange dots) so $166 / 364 = 46 \%$ of the Cavities of the 30 nm ZTC structures would be Exterior and thus relatively easily accessible to the Pd nanoclusters in the colloidal suspension.

A 40 nm ZTC structure would have $40 / 2.5 = 16$ cavities per edge with 16 triangular layers and $1+3+6+10+15+21+28+36+45+55+66+78+91+105+120+136 = 816$ Cavities of which $1+3+6+10+15+21+28+36+45+55+66+78+136 = 500$ would be Interior Cavities and $816 - 500 = 316$ would be Exterior Cavities so $316 / 816 = 39 \%$ of the Cavities of the 40 nm ZTC structures would be Exterior.

Therefore for 30-40 nm ZTC structures only about 1/2 to 1/3 of their Cavities are External so it may be optimal for the number of ZTC Cavities to be 3 x the number of Pd 147-atom nanoclusters.
Can TSC-Jitterbug-Zeolite Fusion produce Abundant Cheap Energy so that Expensive Competition for geologically concentrated Cheap Oil will become unnecessary?

Using total Earth Energy Reserves in Terawatt-years, according to M. Taube, in his book Evolution of Matter and Energy on a Cosmic and Planetary Scale (Springer-Verlag 1985), the number of years that 10^10 people could consume energy at the present USA per capita rate, a consumption rate of about 1,000 Terawatt-years/year, is:

<table>
<thead>
<tr>
<th>Reserves</th>
<th>Duration (years)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oil</td>
<td>850</td>
</tr>
<tr>
<td>Gas</td>
<td>550</td>
</tr>
<tr>
<td>Methane</td>
<td>1,500</td>
</tr>
<tr>
<td>Coal</td>
<td>7,000</td>
</tr>
<tr>
<td>Uranium</td>
<td>1.9 x 10^9</td>
</tr>
<tr>
<td></td>
<td>(1/1000 of Earth supply)</td>
</tr>
<tr>
<td>Thoriurn</td>
<td>7.9 x 10^9</td>
</tr>
<tr>
<td></td>
<td>(1/1000 of Earth supply)</td>
</tr>
<tr>
<td>Deuterium</td>
<td>1.9 x 10^9</td>
</tr>
<tr>
<td></td>
<td>(1/1000 of ocean supply)</td>
</tr>
<tr>
<td>Lithium</td>
<td>1.9 x 10^9</td>
</tr>
<tr>
<td></td>
<td>(source of tritium)</td>
</tr>
</tbody>
</table>

As to solar energy, the total solar energy received by Earth is about 109,000 Terawatt-years/year so that 10^10 people could consume energy at the present USA per capita rate by using about 1% (one percent) of the solar energy received by Earth. This could be done, for example, by building a lot of orbiting solar energy collection dishes and beaming the energy to Earth.

The total geothermal heat flux is about 66 Terawatt-years/year, and the total tidal energy is about 3 Terawatt-years/year, so that those sources would be inadequate to support 10^10 people consuming energy at the present USA per capita rate.
For Everybody on Earth to be Happy, the Abundant Cheap Energy must provide a high Standard of Living (current USA standard) for a lot of people (10 billion), and:

last for a long time (more than decades) - rules out Oil, Gas, Methane, and Coal;

have no serious radioactive waste - rules out Uranium, Thorium, and Tritium (Lithium);

have realistically scalable capital cost - rules out Solar which would require Satellite collectors with area 1% of \( \pi \times 6,000^2 = 1,000,000 \text{ km}^2 = (1,000 \text{ km})^2 \) or cloud-free collectors on Earth surface with the same area. Less than 100% efficiency would require correspondingly larger area of collectors.

That leaves one possible source of Abundant Cheap Energy for 10 billion people:

<table>
<thead>
<tr>
<th>Reserves</th>
<th>Duration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Deuterium</td>
<td>1.9 x 10^9 ( \text{Terawatt-years} )</td>
</tr>
<tr>
<td>(1/1000 of ocean supply)</td>
<td></td>
</tr>
</tbody>
</table>

Since a gram of properly structured Palladium gives TSC-Jitterbug Pd-D Cold Fusion Energy on the scale of Megawatts:

1 milligram of Palladium gives a 1 kiloWatt Machine, useful for “electric motors, tools, machines and heaters” (Wikipedia)
Such small energy machines could use the ZTC Electric Capacitor technology.

1 gram of Palladium gives a 1 MegaWatt = 1340 HorsePower Machine, useful for “large electric motors; large warships such as aircraft carriers, cruisers, and submarines; large server farms or data centers; and some scientific research equipment such as supercolliders, and the output pulses of very large lasers. A large residential or commercial building may use several megawatts in electric power and heat. ... railway... electric locomotives ... typically have a peak power output of 5 or 6 MW, although ... Eurostar ... uses more than 12 MW, while heavy diesel-electric locomotives typically produce/use 3 to 5 MW ...” (Wikipedia)
C-130 aircraft have 4 engines each with 4300 HorsePower (globalsecurity.org) so would need a 4 x 4300 / 1340 = 13 grams of Pd
Such mid-sized energy machines could use, depending on portability and site requirements, either Zeolite Steam or ZTC Electric Capacitor technology.
1 kg of Palladium gives a 1 GigaWatt Machine, useful for “large power plants ... HVDC converters have been built with power ratings up to 2 GW” (Wikipedia). Such machines could use either Zeolite Steam or ZTC Electric Capacitor technology, using HVDC converters up to 2 GW to convert the ZTC Electric Capacitor DC into AC.

1,000 kg = 1 ton of Palladium gives 1 TeraWatt. The total power used by Humans in 2006 was 16 TW. The average lightning strike peaks at 1 TW, but lasts only 30 microseconds. Powerful 20th century lasers produce TW, but only for nanoseconds. (Wikipedia)

1,000 tons of Palladium gives 1 PetaWatt. The Lawrence Livermore Nova laser has power of 1.25 PW in a 5x10^(-13) sec pulse. The total power of sunlight hitting the Earth is about 174 PW. (Wikipedia)

222 tons of Palladium were mined world-wide (based on 2006 and 2007 data, Wikipedia):

- Russia produced 98 tons
- South Africa produced 89 tons
- Canada produced 13 tons
- USA produced 11 tons
- the rest of the world produced 11 tons