

US007604673B2

(12) United States Patent

Kaye et al.

(54) ANNULAR FUEL PROCESSOR AND **METHODS**

- (75) Inventors: Ian W. Kaye, Livermore, CA (US); Arpad Somogyvari, Livermore, CA (US)
- Assignee: UltraCell Corporation, Livermore, CA (73)(US)
- (*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 608 days.
- Appl. No.: 10/877,044 (21)
- (22)Filed: Jun. 25, 2004

Prior Publication Data (65)

US 2005/0011125 A1 Jan. 20, 2005

Related U.S. Application Data

- (60) Provisional application No. 60/482,996, filed on Jun. 27, 2003, provisional application No. 60/483,416, filed on Jun. 27, 2003.
- (51) Int. Cl.

B01J 7/00	(2006.01)
H01M 8/06	(2006.01)
B01J 8/00	(2006.01)
B01J 10/00	(2006.01)
B01J 8/04	(2006.01)
B01J 19/00	(2006.01)

- U.S. Cl. 48/61; 48/127.9; 422/188; (52)422/196; 422/198
- Field of Classification Search 48/127.9, (58)48/61; 422/188, 196, 198 See application file for complete search history.

(56)**References** Cited

U.S. PATENT DOCUMENTS

3,119,671 A	1/1964	Koniewiez et al 422/197
3,541,729 A	11/1970	Dantowitz 48/94

US 7,604,673 B2 (10) Patent No.: (45) Date of Patent:

Oct. 20, 2009

3,578,952 A 5/1971 Boose

(Continued)

FOREIGN PATENT DOCUMENTS

19841993 3/2000

DE

(Continued)

OTHER PUBLICATIONS

U.S. Appl. No. 10/131,846, entitled: Microfluidic Fuel Cell Systems with Embedded Materials and Structures and Method Thereof, by inventors: Morse, filed Apr. 24, 2002.

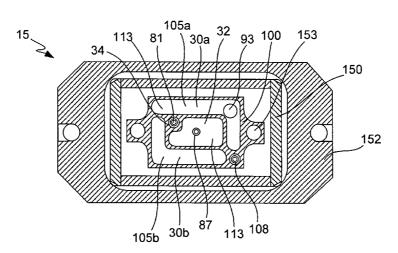
(Continued)

Primary Examiner-Alexa D Neckel Assistant Examiner-Matthew J Merkling (74) Attorney, Agent, or Firm-Beyer Law Group LLP

(57)ABSTRACT

Described herein is a fuel processor that produces hydrogen from a fuel source. The fuel processor comprises a reformer and burner. The reformer includes a catalyst that facilitates the production of hydrogen from the fuel source. Voluminous reformer chamber designs are provided that increase the amount of catalyst that can be used in a reformer and increase hydrogen output for a given fuel processor size. The burner provides heat to the reformer. One or more burners may be configured to surround a reformer on multiple sides to increase thermal transfer to the reformer. Dewars are also described that increase thermal management of a fuel processor and increase burner efficiency. A dewar includes one or more dewar chambers that receive inlet air before a burner receives the air. The dewar is arranged such that air passing through the dewar chamber intercepts heat generated in the burner before the heat escapes the fuel processor.

40 Claims, 13 Drawing Sheets



U.S. PATENT DOCUMENTS

	0.0.1		Decements
3,595,628	Α	7/1971	Connor et al 422/190
4,522,894	Α	6/1985	Hwang et al 429/17
4,909,808	Α	3/1990	Voecks
5,081,095	Α	1/1992	Bedford et al 502/304
5,145,491	Α	9/1992	Schmitt et al.
5,209,906	Α	5/1993	Watkins et al.
5,525,436	Α	6/1996	Savinell et al 429/30
5,534,328	Α	7/1996	Ashmead et al 428/166
5,601,938	Α	2/1997	Mayer et al 429/40
5,611,214	Α	3/1997	Wegeng et al 62/498
5,641,585	Α	6/1997	Lessing et al 429/26
5,716,727	Α	2/1998	Savinell et al 429/33
5,733,347	Α	3/1998	Lesieur 422/129
5,789,093	Α	8/1998	Malhi 429/34
5,811,062	Α	9/1998	Wegeng et al 422/129
5,858,314	A *	1/1999	Hsu et al 422/211
5,961,930	Α	10/1999	Chatterjee et al 422/130
5,961,932	Α	10/1999	Ghosh et al 422/193
6,077,620	Α	6/2000	Pettit 429/26
6,080,501	Α	6/2000	Kelley et al 429/31
6,193,501	B1	2/2001	Masel et al 431/170
6,200,536	B1	3/2001	Tonkovich et al 422/177
6,232,005	B1	5/2001	Pettit
6,245,214	B1	6/2001	Rehg et al 205/764
6,268,077	B1	7/2001	Kelley et al 429/33
6,312,846	B1	11/2001	Marsh 429/30
6,406,808	B1	6/2002	Pratt et al 429/26
6,415,860	B1	7/2002	Kelly et al 165/148
6,423,434	B1	7/2002	Pratt et al 429/13
6,460,733	B2	10/2002	Acker et al 222/94
6,465,119	B1	10/2002	Koripella et al 429/32
6,470,569	B1	10/2002	Lippert et al 29/890
6,537,352	B2 *	3/2003	Edlund et al 96/4
6,537,506	B1	3/2003	Schwalbe et al 422/130
6,541,676	B1	4/2003	Franz et al 585/250
6,569,553	B1	5/2003	Koripella et al 429/20
6,638,654	B2	10/2003	Jankowski et al 429/26
6,645,443	B1	11/2003	Vogel et al.
6,673,130	B2	1/2004	Jankowski et al 29/623.5
6,753,036	B2	6/2004	Jankowski et al 427/189
6,770,106	B1 *	8/2004	Okamoto et al 48/127.9
6,810,899		11/2004	Franz et al 137/79
6,821,666	B2	11/2004	Morse et al 429/34 Allen et al.
6,887,587	B2 D2*	5/2005 8/2005	Wangerow et al 423/652
6,932,958	B2 * B2		Arana et al 429/17
6,939,632 7,014,835	B2 *	9/2005 3/2006	Mathias et al 423/652
7,056,477	B1	6/2006	Schwalbe et al
7,115,148	B2	10/2006	Wheeldon et al.
7,214,251	B2 *	5/2007	Oh et al
2001/0029735	Al	10/2001	Miura et al.
2001/0029974	Al	10/2001	Cohen et al 136/201
2001/0031592	Al	10/2001	Schulz
2002/0012825	Al	1/2002	Sasahara et al 429/30
2002/0045082	A1	4/2002	Marsh 429/30
2002/0076599	Al	6/2002	Neutzler et al 429/38
2002/0081468	Al	6/2002	Shioya 429/19
2002/0094462	A1	7/2002	Shioya et al 429/19
2002/0098119	A1	7/2002	Goodman 422/82.01
2002/0106540	A1	8/2002	Shioya 429/19
2002/0127141	A1	9/2002	Acker 422/40
2002/0131915	A1	9/2002	Shore et al 422/177
2002/0132156	A1	9/2002	Ruhl et al 429/40
2002/0147107	A1	10/2002	Abdo et al 502/232
2002/0150804	A1	10/2002	Srinivasan et al 429/32
2002/0155335	A1	10/2002	Kearl 429/30
2002/0192537	A1	12/2002	Ren 429/44
2003/0006668	A1	1/2003	Lal et al 310/301
2003/0027022		2/2003	Arana et al 429/17
2003/0031910	A1	2/2003	Satou et al 429/30
2003/0031913	A1	2/2003	Pavio et al 429/34

2003/0057199	A1	3/2003	Villa et al 219/521
2003/0077490	A1	4/2003	Reinke et al.
2003/0091502	A1	5/2003	Holladay et al 423/652
2003/0129464	A1	7/2003	Becerra et al 429/25
2003/0159354	A1	8/2003	Edlund et al.
2003/0190508	A1	10/2003	Takeyama et al 429/20
2003/0194363	A1	10/2003	Koripella 422/222
2003/0198592	A1*	10/2003	Allison et al 423/651
2004/0009381	A1	1/2004	Sakai et al 429/23
2004/0025784	A1	2/2004	Kawamura et al 117/200
2004/0028602	A1	2/2004	Franz et al 423/648.1
2004/0031592	A1	2/2004	Mathias et al.
2004/0040281	A1	3/2004	Yamaguchi et al.
2004/0043263	A1	3/2004	Takeyama et al 429/19
2004/0062961	A1	4/2004	Sato et al 429/19
2004/0065013	A1	4/2004	DeVries 48/198.2
2004/0105789	A1	6/2004	Yamamoto 422/130
2004/0228211	A1	11/2004	Koripella et al 366/341
2005/0008911	A1	1/2005	Kaye
2005/0011125	A1	1/2005	Kaye et al.
2005/0115889	A1	6/2005	Schaevitz et al 210/490
2006/0107594	A1	5/2006	Kuwaba
2006/0127719	A1	6/2006	Brantley et al.
2006/0134470	A1	6/2006	Kaye et al.
2008/0118796	A1	5/2008	Brantley et al.

FOREIGN PATENT DOCUMENTS

06-310166	Α	11/1994
2003048701		2/2003
2003290649		10/2003
2003290653		10/2003
2003340273		12/2003
2004057920		2/2004
WO 00/45457		8/2000
WO 0147800	A1	* 7/2001
WO02/19452		3/2002
WO 02/059993		8/2002
WO 02/093665		11/2002
WO 02/103832		12/2002
WO 02/103878		12/2002
WO03/082460		10/2003
WO03/083984		10/2003
WO 2004/030805		4/2004
WO2004/037406		5/2004
	2003048701 2003290649 2003290653 2003340273 2004057920 WO 00/45457 WO 0147800 WO02/19452 WO 02/059993 WO 02/099365 WO 02/103832 WO 02/103878 WO03/082460 WO03/083984 WO 2004/030805	2003290649 2003290653 2003340273 2004057920 WO 00/45457 WO 0147800 A1 WO02/19452 WO 02/059993 WO 02/09365 WO 02/103878 WO 02/103878 WO03/082460 WO03/083984 WO 2004/030805

OTHER PUBLICATIONS

U.S. Appl. No. 10/261,353, entitled: Bonded Polyimide Fuel Cell Package and Method Thereof, by inventors: Morse, filed Sep. 30, 2002.

U.S. Appl. No. 10/371,876, entitled: Metal Hydride Fuel Storage and Method Thereof, by inventors: Morse, filed Feb. 21, 2003.

U.S. Appl. No. 10/612,177, entitled: Vapor-deposited Porous Films for Energy Conversion, by inventors: Jankowski, filed Jul. 1, 2003. U.S. Appl. No. 10/637,914, entitled: Solid Oxide MEMS-based Fuel

Cells, by inventors: Jankowski, filed Aug. 8, 2003.

U.S. Appl. No. 10/637,915, entitled: Solid Polymer MEMS-based Fuel Cells, by inventors: Jankowski, filed Aug. 8, 2003.

U.S. Appl. No. 10/783,230, entitled: Method for Fabrication of Electrodes, by inventors: Jankowski, filed Feb. 19, 2004.

U.S. Appl. No. 10/853,859, entitled: Microfluidic Systems with Embedded Materials and Structures and Method Thereof, by inventors: Morse, filed May 25, 2004.

U.S. Appl. No. 10/877,769, entitled "Micro Fuel Cell System Start Up and Shut Down Systems and Methods", by inventors: Kaye, filed Jun. 25, 2004.

U.S. Appl. No. 10,877,826, entitled "Fuel Preheat in Fuel Cells and Portable Electronics", by inventors: Kaye, filed Jun. 25, 2004.

U.S. Appl. No. 10/877,771, entitled "Efficient Micro Fuel Cell Systems and Methods", by inventors: Kaye, filed Jun. 25, 2004.

U.S. Appl. No. 10/007,412, entitled "Chemical Microreactor and Method Thereof", by inventors: Morse, filed Dec. 5, 2001.

S. Ahmed et al., "Catalytic Partial Oxidation Reforming of Hydrocarbon Fuels", Nov. 16-19, 1998, 1998 Fuel Cell Seminar, Palm Springs, CA.

A.R. Boccaccini et al., "Electrophoretic Deposition of Nanoceramic Particles onto Electrically Conducting Fibre Fabrics", Sep. 21-24, 1998, 43rd International Scientific Colloquium, Technical University of Ilmenau.

J. Bostaph et al., "1 W Direct Methanol Fuel Cell System as a Desktop Charger", Oct. 14, 2002, Motorola Labs, Tempe, AZ.

K. Brooks et al., "Microchannel Fuel Processing, Fuel Cells for Transportation/Fuels for Fuel Cells", May 6-10, 2002, 2002 Annual Program/Lab R&D Review, Pacific Northwest National Laboratory. M.J. Castaldi et al., "A Compact, Lightweight, Fast-Response Preferential Oxidation Reactor for PEM Automotive Fuel Cell Applications", Sep. 6, 2002, Precision Combustion, Inc., North Haven, CT. S. Ehrenberg et al., "One Piece Bi-Polar (OPB) Plate with Cold Plate Cooling", Dec. 13, 2002, Session PEM R&D II (2A), Dais Analytic—Rogers.

T.M. Floyd et al., "Liquid-Phase and Multi-Phase Microreactors for Chemical Synthesis", Jun. 1999, Massachusetts Institute of Technology, Cambridge, MA.

A.J. Franz et al., "High Temperature Gas Phase Catalytic and Membrane Reactors", Jun. 1999, Massachusetts Institute of Technology, Cambridge, MA.

J.D. Holladay et al., "Miniature Fuel Processors for Portable Fuel Cell Power Supplies", Nov. 26, 2002, Battelle Pacific Northwest Division, Richland, WA.

S.W. Janson et al., "MEMS, Microengineering and Aerospace Systems", 1999, The American Institute of Aeronautics and Astronautics, Inc.

J. Kaschmitter et al., "Micro-Fabricated Methanol/Water Reformers for Small PEM Fuel Cell Systems", Jul. 21-24, 2003, 8th Electrochemical Power Sources R&D Symposium, Portsmouth, VA.

K. Keegan et al., "Analysis for a Planar Solid Oxide Fuel Cell Based Automotive Auxiliary Power Unit", Mar. 4-7, 2002, SAE 2002 World Congress, Detroit, MI.

K. Kempa et al., "Photonic Crystals Based on Periodic Arrays of Aligned Carbon Nanotubes", Oct. 3, 2002, Nano Letters 2003, vol. 3. No. 1, 13-18.

R. Kumar et al., "Solid Oxide Fuel Cell Research at Argonne National Laboratory", Mar. 29-30, 2001, 2nd Solid Sate Energy Conversion Alliance Workshop, Arlington, VA.

S.H. Lee et al., "Removal of Carbon Monoxide from Reformate for Polymer Electrolyte Fuel Cell Application", Nov. 16-19, 1998, 1998 Fuel Cell Seminar, Palm Springs, CA.

Q. Li et al., "The CO Poisoning Effect in PEMFCs Operational at Temperatures up to 200° C.", 2003, Journal of The Electrochemical Society, 150 (12) A1599-A1605.

D. Myers et al., "Alternative Water-Gas Shift Catalysts", Jun. 7-8, 2000, 2000 Annual National Laboratory R&D Meeting, DOE Fuel Cells for Transportation Program, Argonne National Laboratory.

D.R. Palo et al., "Development of a Soldier-Portable Fuel Cell Power System, Part I: A Bread-Board Methanol Fuel Processor", 2002, Journal of Power Sources 108 (2002) 28-34.

A. Pattekar et al., "A Microreactor for In-situ Hydrogen Production by Catalytic Methanol Reforming", May 27-30, 2001, Proceedings of the $5'^{th}$ International Conference on Microreaction Technology.

A. Pattekar et al., "Novel Microfluidic Interconnectors for High Temperature and Pressure Applications", 2003, Journal of Micromechanics and Microengineering, 13, 337-345. D. Prater et al., "Systematic Examination of a Direct Methanol-Hydrogen Peroxide Fuel Cell", Sep. 22, 2001, Swift Enterprises, Ltd., Lafayette, IN.

W. Ruettinger et al., "A New Generation of Water Gas Shift Catalysts for Fuel Cell Applications", 2003, Journal of Power Sources, 118, 61-65.

O. Savadogo et al., Hydrogen/Oxygen Polymer Electrolyte Membrane Fuel Cell (PEMFC) Based on Acid-Doped Polybenzimidazole (PBI), 2000, Journal of New Materials for Electrochemical Systems, 3, 345-349.

R.F. Savinell et al., "High Temperature Polymer Electrolyte for PEM Fuel Cells", Sep. 4, 2002, Department of Chemical Engineering, Case Western Reserve University.

R. Srinivasan et al., "Micromachined Reactors for Catalytic Partial Oxidation Reactions", Nov. 1997, AIChe Journal, vol. 43, No. 11, 3059-3069.

S. Swartz et al., "Ceria-Based Water-Gas-Shift Catalysts", Aug. 1, 2003, NexTech Materials, Ltd., Wolrthington, OH.

S. Tasic et al., "Multilayer Ceramic Processing of Microreactor Systems", Oct. 14, 2002, Motorola Labs, Tempe, AZ.

V. Tomašić et al., "Development of the Structured Catalysts for the Exhaust Gas Treatment", 2001, Chem. Biochem. Eng. Q. 15 (3), 109-115.

Tiax LLC, "Advanced Hydrogen Storage: A System's Perspective and Some Thoughts on Fundamentals", Aug. 14-15, 2002, Presentation for DOE Workshop on Hydrogen Storage, Cambridge, MA.

Wan et al., "Catalyst Preparation: Catalytic Converter", Feb. 19, 2003, www.insightcentral.net/encatalytic.html.

J. Zalc et al., "Are Noble Metal-Based Water-Gas Shift Catalysts Practical for Automotive Fuel Processing?", 2002, Journal of Catalysis, 206, 169-171.

J. Zizelman et al., "Solid-Oxide Fuel Cell Auxiliary Power Unit: A Paradigm Shift in Electric Supply for Transportation", undated, Delphi Automotive Systems.

"Methanol-Powered Laptops—Cleared for Take-Off", www.silicon. com, Oct. 7, 2002.

Melissa Funk, "Methanol Fuel Quality Specification Study for Proton Exchange Membrane Fuel Cells, Final Report", XCELLSIS, Feb. 2002, 65 pages.

Dr. Detlef zur Megede et al., "MFCA Research Document, Complete", Methanol Fuel Cell Alliance, Sep. 2000, 242 pages.

Office Action dated Jan. 25, 2008 in U.S. Appl. No. 10/877,804.

Office Action dated Apr. 16, 2008 in U.S. Appl. No. 11/313,252. Office Action dated Jun. 3, 2008 from Indian Patent Application No. 16/KOLNP/2006.

Office Action dated Nov. 6, 2008 in U.S. Appl. No. 11/830,177.

Office Action dated Oct. 15, 2008 from U.S. Appl. No. 11/313,252. Notice of Allowance dated Sep. 25, 2008 from U.S. Appl. No. 10/877,804.

Conduction General Theory, Efunda, Feb. 2002.

Office Action dated Jun. 10, 2009 in U.S. Appl. No. 11/830,181.

Office Action dated Apr. 24, 2009 in U.S. Appl. No. 11/830,177.

Office Action dated Apr. 29, 2009 from U.S. Appl. No. 11/772,689.

Office Action dated Mar. 30, 2009 in U.S. Appl. No. 11/313,252.

Office Action dated Dec. 22, 2008 from U.S. Appl. No. 11/772,689. International Search Report dated Oct. 22, 2008 from PCT Application No. PCT/US07/17790.

Written Opinion of International Search Authority dated Oct. 22, 2008 from PCT Application No. PCT/US07/17790.

Office Action dated Dec. 24, 2008 from U.S. Appl. No. 11/835,747.

* cited by examiner

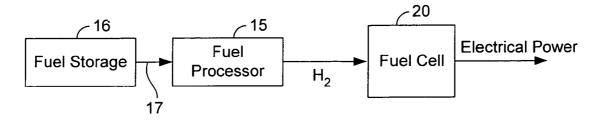
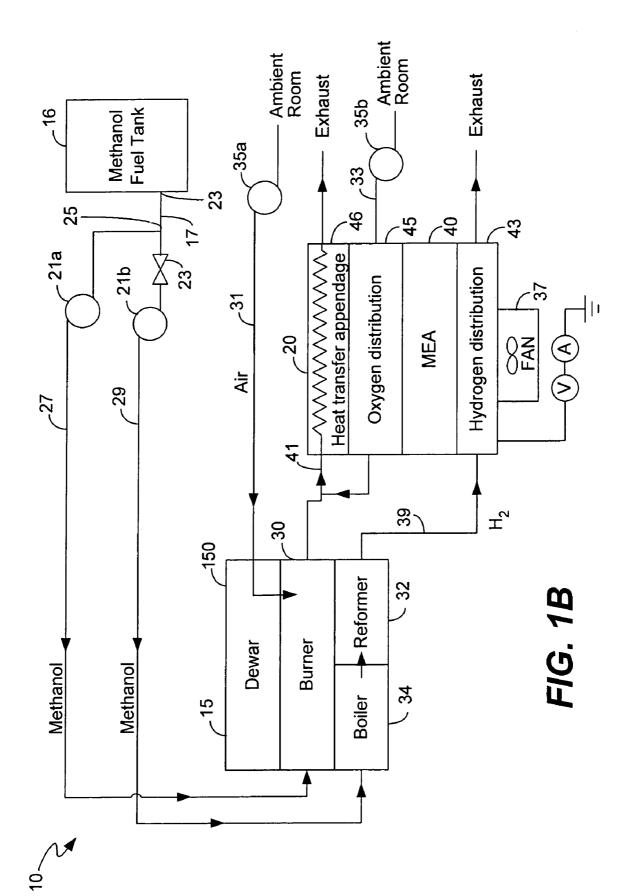
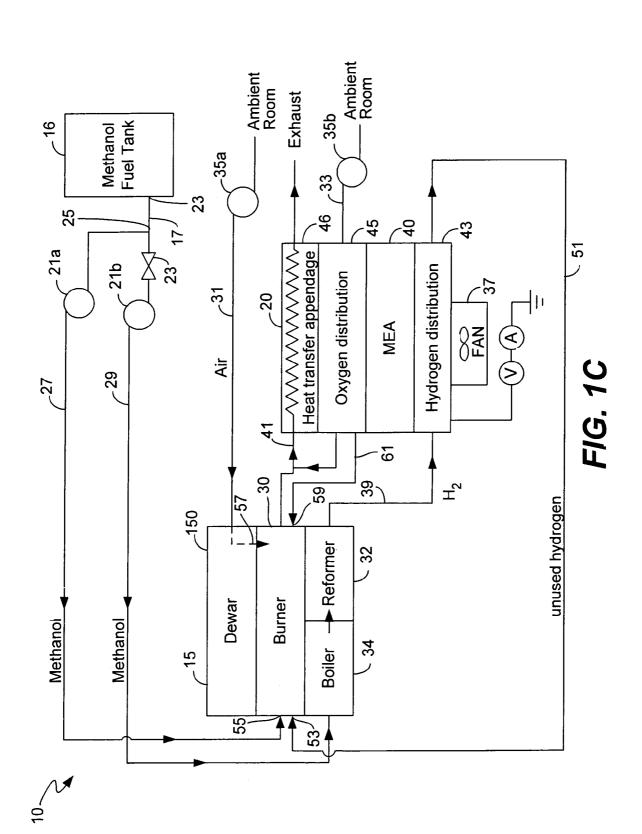


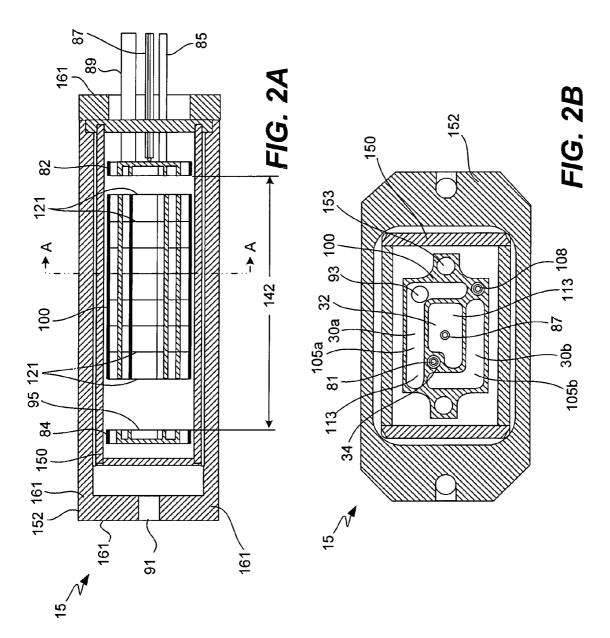
FIG. 1A



U.S. Patent



Sheet 3 of 13



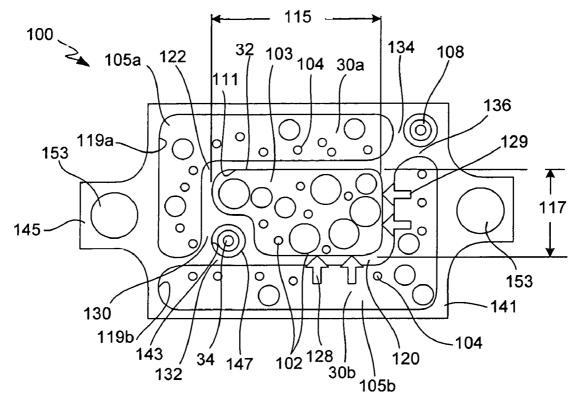
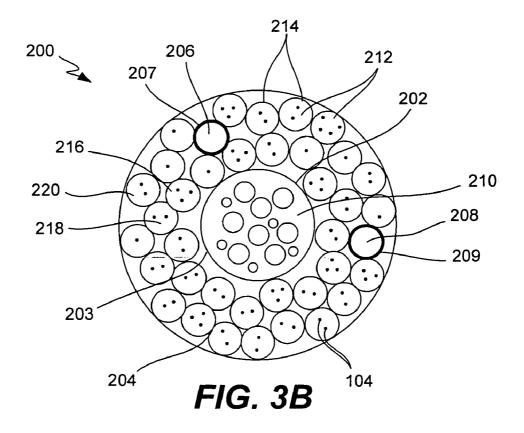


FIG. 3A



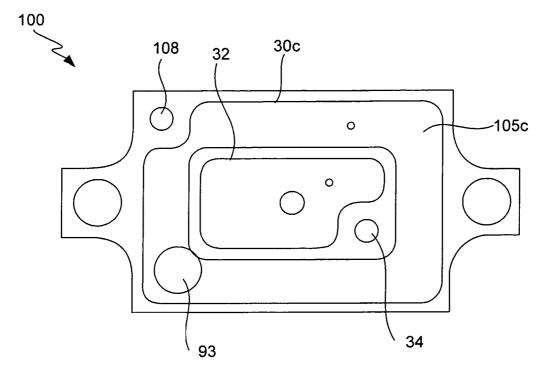
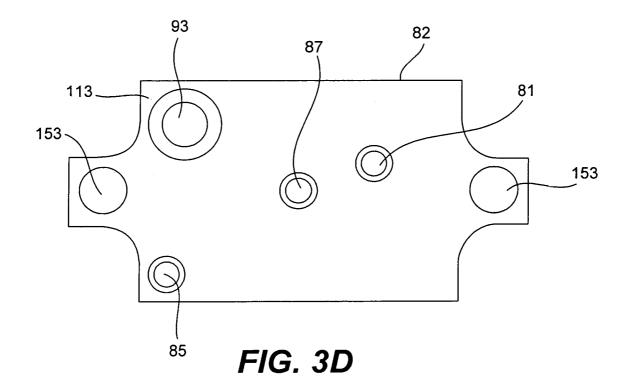


FIG. 3C



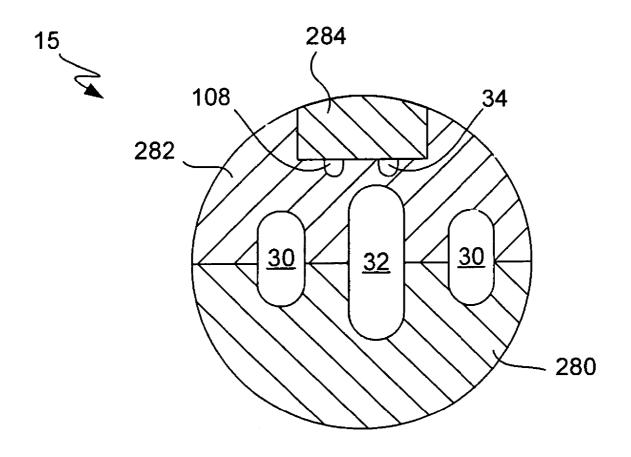
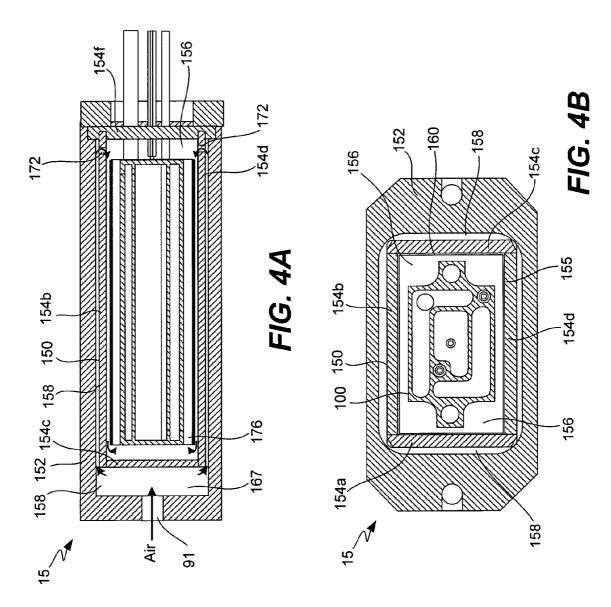


FIG. 3E



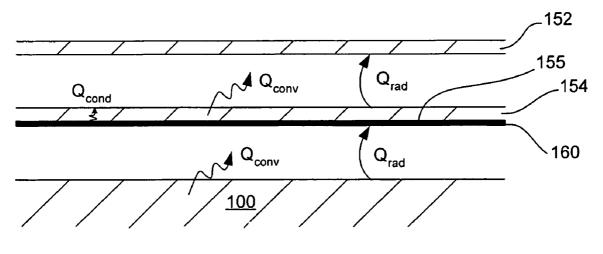


FIG. 4C

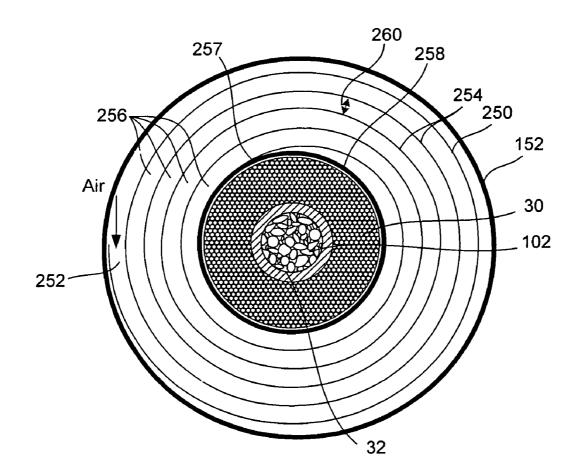
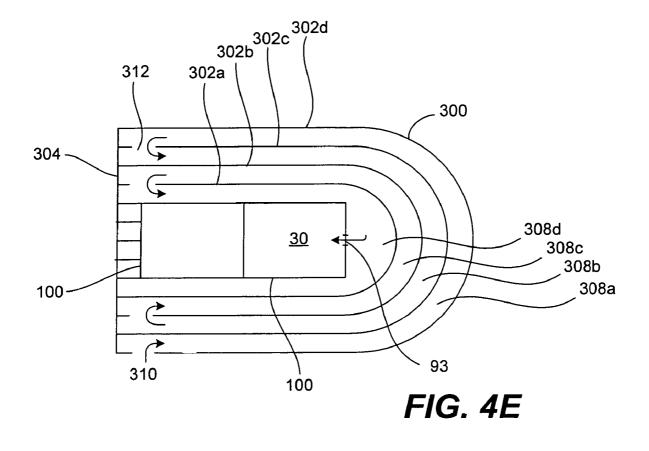
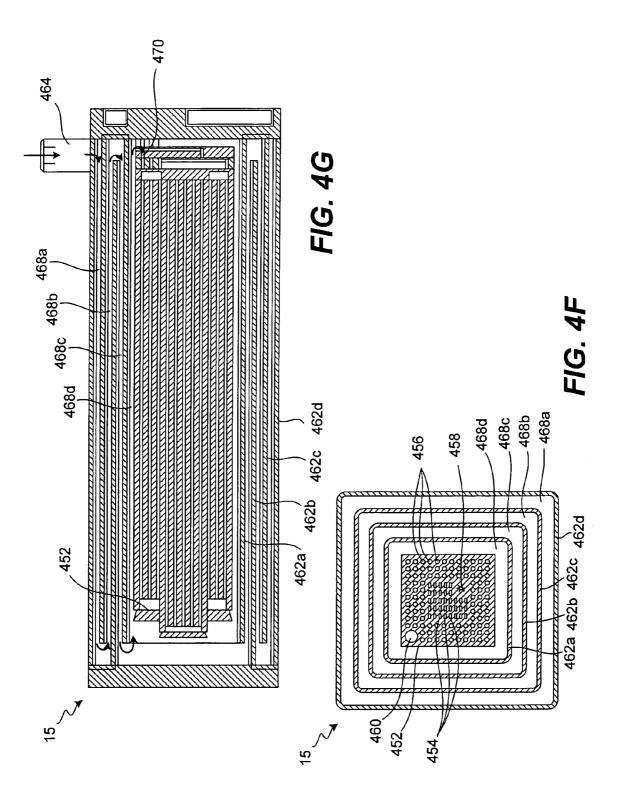
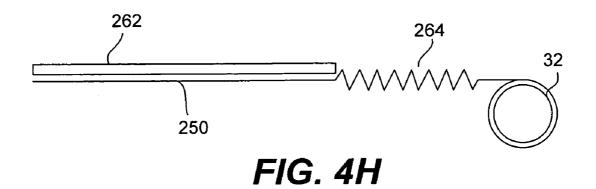
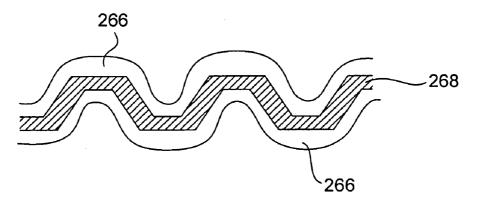


FIG. 4D











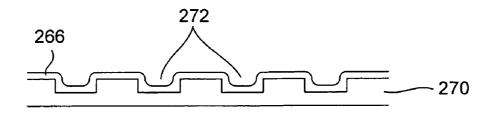
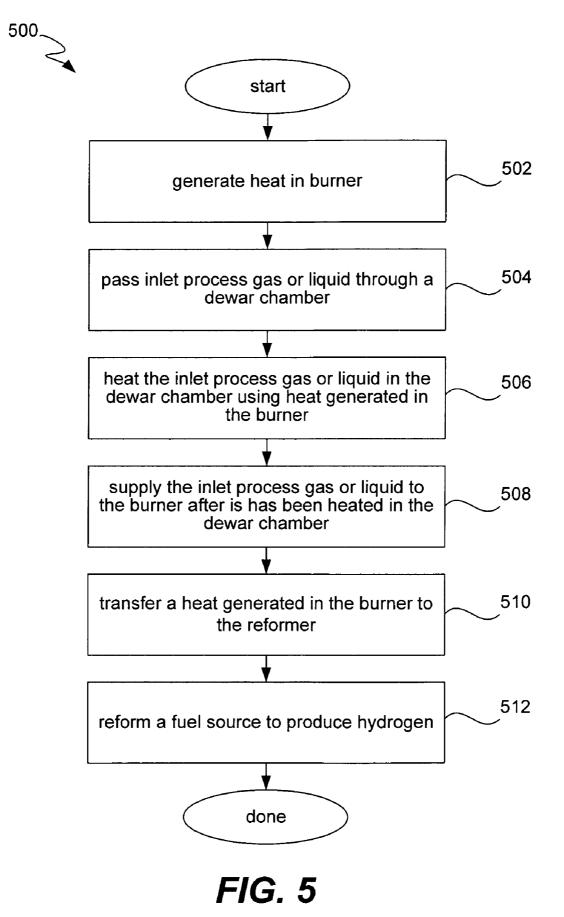


FIG. 4J



ANNULAR FUEL PROCESSOR AND METHODS

CROSS-REFERENCE TO RELATED APPLICATIONS

This application claims priority under 35 U.S.C. §119(e) from U.S. Provisional Patent Application No. 60/482,996 filed Jun. 27, 2003 entitled "Fuel cell system startup procedure and self-heating apparatus", which is incorporated by 10 reference for all purposes; and also claims priority under 35 U.S.C. §119(e) from U.S. Provisional Patent Application No. 60/483,416 filed Jun. 27, 2003 entitled "Fuel Preheat in Portable Electronics Powered by Fuel Cells", which is incorporated by reference for all purposes. 15

BACKGROUND OF THE INVENTION

The present invention relates to fuel cell technology. In particular, the invention relates to fuel processors that gener- ²⁰ ate hydrogen and are suitable for use in portable applications.

A fuel cell electrochemically combines hydrogen and oxygen to produce electricity. The ambient air readily supplies oxygen. Hydrogen provision, however, calls for a working supply. Gaseous hydrogen has a low energy density that 25 reduces its practicality as a portable fuel. Liquid hydrogen, which has a suitable energy density, must be stored at extremely low temperatures and high pressures, making storing and transporting liquid hydrogen burdensome.

A reformed hydrogen supply processes a fuel source to 30 produce hydrogen. The fuel source acts as a hydrogen carrier. Currently available hydrocarbon fuel sources include methanol, ethanol, gasoline, propane and natural gas. Liquid hydrocarbon fuel sources offer high energy densities and the ability to be readily stored and transported. A fuel processor reforms 35 the hydrocarbon fuel source and to produce hydrogen.

Fuel cell evolution so far has concentrated on large-scale applications such as industrial size generators for electrical power back-up. Consumer electronics devices and other portable electrical power applications currently rely on lithium 40 ion and similar battery technologies. Fuel processors for portable applications such as electronics would be desirable but are not yet commercially available. In addition, techniques that reduce fuel processor size or increase fuel processor efficiency would be highly beneficial. 45

SUMMARY OF THE INVENTION

The present invention relates to a fuel processor that produces hydrogen from a fuel source. The fuel processor comprises a reformer and burner. The reformer includes a catalyst that facilitates the production of hydrogen from the fuel source. Voluminous reformer chamber designs are provided that increase the amount of catalyst that can be used in a reformer and increase hydrogen output for a given fuel pro-55 cessor size. The burner provides heat to the reformer. One or more burners may be configured to surround a reformer on multiple sides to increase thermal transfer to the reformer.

Dewars are also described that improve thermal management of a fuel processor by reducing heat loss and increasing 60 burner efficiency. A dewar includes one or more dewar chambers that receive inlet process gases or liquids before a reactor receives them. The dewar is arranged such that inlet process gases or liquids passing through the dewar chamber intercepts heat generated in the burner before the heat escapes the fuel 65 processor. Passing inlet process gases or liquids through a dewar chamber in this manner performs three functions: a)

active cooling of dissipation of heat generated in burner before is reaches outer portions of the fuel processor, and b) heating of the air before receipt by the burner, and c) absorbing and recycling heat back into the burner increasing burner efficiency. When the burner relies on catalytic combustion to produce heat, heat generated in the burner warms cool process gases or liquids in the burner according to the temperature of the process gases or liquids. This steals heat from the reformer, reduces heating efficiency of a burner and typically results in greater consumption of the fuel source. The dewar thus pre-heats the incoming process gases or liquids before burner arrival so the burner passes less heat to the process gases or liquids that would otherwise transfer to the reformer.

In one aspect, the present invention relates to a fuel processor for producing hydrogen from a fuel source. The fuel processor comprises a reformer configured to receive the fuel source, configured to output hydrogen, and including a catalyst that facilitates the production of hydrogen. The fuel processor also comprises a boiler configured to heat the fuel 20 source before the reformer receives the fuel source. The fuel processor further comprises at least one burner configured to provide heat to the reformer and disposed annularly about the reformer. The fuel processor may also comprise a boiler that heats the burner liquid fuel feed.

In another aspect, the present invention relates to a fuel processor for producing hydrogen from a fuel source. The fuel processor comprises a reformer configured to receive the fuel source, configured to output hydrogen, including a catalyst that facilitates the production of hydrogen. The reformer also includes a reformer chamber having a volume greater than about 0.1 cubic centimeters and less than about 50 cubic centimeters and is characterized by a cross sectional width and a cross sectional height that is greater than one-third the cross sectional width. The fuel processor also comprises a boiler configured to heat the fuel source before the reformer receives the fuel source. The fuel processor further comprises at least one burner configured to provide heat to the reformer.

In yet another aspect, the present invention relates to a fuel processor for producing hydrogen from a fuel source. The fuel processor comprises a reformer configured to receive the hydrogen fuel source, configured to output hydrogen, and including a catalyst that facilitates the production of hydrogen. The fuel processor also comprises a burner that is configured to provide heat to the reformer. The fuel processor further comprises a dewar that at least partially contains the reformer and the burner and includes a set of dewar walls that form a dewar chamber that is configured to receive an inlet process gas or liquid before the burner receives the inlet process gas or liquid. The fuel processor additionally comprises a housing including a set of housing walls that at least partially contain the dewar and provide external mechanical protection for the reformer and the burner.

In still another aspect, the present invention relates to a method for managing heat in a fuel processor. The fuel processor comprises a burner, a reformer and a dewar that at least partially contains the burner. The method comprises generating heat in the burner. The method also comprises passing an inlet process gas or liquid through a dewar chamber. The method further comprises heating the inlet process gas or liquid in the dewar chamber using heat generated in the burner.

In another aspect, the present invention relates to a method for generating hydrogen in a fuel processor. The fuel processor comprises a burner, a reformer and a dewar that at least partially contains the burner and reformer. The method comprises generating heat in the burner. The method also comprises passing an inlet process gas or liquid through a dewar

60

chamber. The method further comprises heating the inlet process gas or liquid in the dewar chamber using heat generated in the burner. The method additionally comprises supplying the inlet process gas or liquid to the burner after it has been heated in the dewar chamber. The method also comprises transferring heat generated in the burner to the reformer. The method further comprises reforming a fuel source to produce hydrogen.

These and other features and advantages of the present invention will be described in the following description of the 10 invention and associated figures.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. **1**A illustrates a fuel cell system for producing elec-¹⁵ trical energy in accordance with one embodiment of the present invention.

FIG. 1B illustrates schematic operation for the fuel cell system of FIG. 1A in accordance with a specific embodiment of the present invention. 20

FIG. 1C illustrates an embodiment of the fuel cell system of FIG. 1A that routes hydrogen from an anode exhaust of the fuel cell back to a burner in the fuel processor.

FIG. **2**A illustrates a cross-sectional side view of a fuel processor used in the fuel cell system of FIG. **1**A in accordance with one embodiment of the present invention.

FIG. **2**B illustrates a cross-sectional front view of the fuel processor used in the fuel cell system of FIG. **1**A taken through a mid-plane of fuel processor.

FIG. **3**A illustrates a cross-sectional front view of a monolithic structure employed in the fuel processor of FIG. **2**A in accordance with one embodiment of the present invention.

FIG. **3**B illustrates a cross-sectional layout of a tubular design for use in a fuel processor in accordance with another $_{35}$ embodiment of the present invention.

FIG. **3**C illustrates a cross-sectional front view of a monolithic structure in a fuel processor that comprises a single burner having an 'O-shape' that completely surrounds a reformer chamber in accordance with one embodiment of the $_{40}$ present invention.

FIG. **3**D illustrates an outside view of an end plate used in the fuel processor of FIG. **2**A.

FIG. **3**E illustrates a fuel processor **15** in accordance with another embodiment of the present invention.

FIG. **4**A illustrates a side cross-sectional view of the fuel processor of FIG. **2**A and movement of air created by a dewar in accordance with one embodiment of the present invention.

FIG. **4**B illustrates a front cross-sectional view of the fuel processor of FIG. **2**A and demonstrates thermal management ⁵⁰ benefits gained by the dewar.

FIG. **4**C shows a thermal diagram of the heat path produced by a dewar wall used in the fuel processor of FIG. **2**A.

FIG. 4D illustrates a cross sectional view of a fuel processor that increases the convective path that air flows over a dewar wall in accordance with another embodiment of the present invention.

FIG. **4**E illustrates a dewar in accordance with another embodiment of the present invention.

FIGS. 4F and 4G illustrate a cross section of a fuel processor including a monolithic structure and multipass dewar in accordance with another embodiment of the present invention.

FIG. **4**H illustrates spiral dewar in an unrolled form during 65 initial construction in accordance with another embodiment of the present invention.

4

FIGS. 4I and 4J illustrate wash coatings on a wall of a burner in accordance with two embodiments of the present invention.

FIG. **5** illustrates a process flow for generating hydrogen in a fuel processor in accordance with one embodiment of the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is described in detail with reference to a few preferred embodiments as illustrated in the accompanying drawings. In the following description, numerous specific details are set forth in order to provide a thorough understanding of the present invention. It will be apparent, however, to one skilled in the art, that the present invention may be practiced without some or all of these specific details. In other instances, well known process steps and/or structures have not been described in detail in order to not unnecessarily obscure the present invention.

FIG. 1A illustrates a fuel cell system 10 for producing electrical energy in accordance with one embodiment of the present invention. Fuel cell system 10 comprises storage device 16, fuel processor 15 and fuel cell 20.

A 'reformed' hydrogen supply processes a fuel source to produce hydrogen. As shown, the reformed hydrogen supply comprises a fuel processor 15 and a fuel source storage device 16. Storage device 16 stores fuel source 17, and may include a portable and/or disposable fuel cartridge. A disposable cartridge offers instant recharging to a consumer. In one embodiment, the cartridge includes a collapsible bladder within a hard plastic dispenser case. A separate fuel pump typically controls fuel source 17 flow from storage device 16. If system 10 is load following, then a control system meters fuel source 17 to deliver fuel source 17 to processor 15 at a flow rate determined by the required power level output of fuel cell 20.

Fuel source 17 acts as a carrier for hydrogen and can be processed to separate hydrogen. Fuel source 17 may include any hydrogen bearing fuel stream, hydrocarbon fuel or other hydrogen fuel source such as ammonia. Currently available hydrocarbon fuel sources 17 suitable for use with the present invention include gasoline, C1 to C4 hydrocarbons, their oxygenated analogues and/or their combinations, for example. Several hydrocarbon and ammonia products may also produce a suitable fuel source 17. Liquid fuel sources 17 offer high energy densities and the ability to be readily stored and shipped. Storage device 16 may contain a fuel mixture. When the fuel processor 15 comprises a steam reformer, storage device 16 may contain a fuel mixture of a hydrocarbon fuel source and water. Hydrocarbon fuel source/water fuel mixtures are frequently represented as a percentage fuel source in water. In one embodiment, fuel source 17 comprises methanol or ethanol concentrations in water in the range of 1%-99.9%. Other liquid fuels such as butane, propane, gasoline, military grade "JP8" etc. may also be contained in storage device 16 with concentrations in water from 5-100%. In a specific embodiment, fuel source 17 comprises 67% methanol by volume.

Fuel processor **15** processes the hydrocarbon fuel source **17** and outputs hydrogen. A hydrocarbon fuel processor **15** heats and processes a hydrocarbon fuel source **17** in the presence of a catalyst to produce hydrogen. Fuel processor **15** comprises a reformer, which is a catalytic device that converts a liquid or gaseous hydrocarbon fuel source **17** into hydrogen and carbon dioxide. As the term is used herein, reforming refers to the process of producing hydrogen from a fuel source. Fuel processor **15** may output either pure hydrogen or a hydrogen bearing gas stream. Fuel processor **15** is described in further detail below.

Fuel cell **20** electrochemically converts hydrogen and oxygen to water, generating electricity and heat in the process. 5 Ambient air commonly supplies oxygen for fuel cell **20**. A pure or direct oxygen source may also be used for oxygen supply. The water often forms as a vapor, depending on the temperature of fuel cell **20** components. The electrochemical reaction also produces carbon dioxide as a byproduct for 10 many fuel cells.

In one embodiment, fuel cell 20 is a low volume polymer electrolyte membrane (PEM) fuel cell suitable for use with portable applications such as consumer electronics. A polymer electrolyte membrane fuel cell comprises a membrane 15 electrode assembly 40 that carries out the electrical energy generating electrochemical reaction. The membrane electrode assembly 40 includes a hydrogen catalyst, an oxygen catalyst and an ion conductive membrane that a) selectively conducts protons and b) electrically isolates the hydrogen 20 catalyst from the oxygen catalyst. A hydrogen gas distribution layer contains the hydrogen catalyst and allows the diffusion of hydrogen therethrough. An oxygen gas distribution layer contains the oxygen catalyst and allows the diffusion of oxygen and hydrogen protons therethrough. The ion conduc- 25 tive membrane separates the hydrogen and oxygen gas distribution layers. In chemical terms, the anode comprises the hydrogen gas distribution layer and hydrogen catalyst, while the cathode comprises the oxygen gas distribution layer and oxygen catalyst.

A PEM fuel cell often includes a fuel cell stack having a set of bi-polar plates. A membrane electrode assembly is disposed between two bi-polar plates. Hydrogen distribution **43** occurs via a channel field on one plate while oxygen distribution **45** occurs via a channel field on a second facing plate. 35 Specifically, a first channel field distributes hydrogen to the hydrogen gas distribution layer, while a second channel field distributes oxygen to the oxygen gas distribution layer. The 'term 'bi-polar' refers electrically to a bi-polar plate (whether comprised of one plate or two plates) sandwiched between 40 two membrane electrode assembly layers. In this case, the bi-polar plate acts as both a negative terminal for one adjacent membrane electrode assembly and a positive terminal for a second adjacent membrane electrode assembly arranged on the opposite face of the bi-polar plate. 45

In electrical terms, the anode includes the hydrogen gas distribution layer, hydrogen catalyst and bi-polar plate. The anode acts as the negative electrode for fuel cell **20** and conducts electrons that are freed from hydrogen molecules so that they can be used externally, e.g., to power an external 50 circuit. In a fuel cell stack, the bi-polar plates are connected in series to add the potential gained in each layer of the stack. In electrical terms, the cathode includes the oxygen gas distribution layer, oxygen catalyst and bi-polar plate. The cathode represents the positive electrode for fuel cell **20** and conducts 55 the electrons back from the external electrical circuit to the oxygen catalyst, where they can recombine with hydrogen ions and oxygen to form water.

The hydrogen catalyst separates the hydrogen into protons and electrons. The ion conductive membrane blocks the electrons, and electrically isolates the chemical anode (hydrogen gas distribution layer and hydrogen catalyst) from the chemical cathode. The ion conductive membrane also selectively conducts positively charged ions. Electrically, the anode conducts electrons to a load (electricity is produced) or battery (energy is stored). Meanwhile, protons move through the ion conductive membrane, to combine with oxygen. The protons

and used electrons subsequently meet on the cathode side, and combine with oxygen to form water. The oxygen catalyst in the oxygen gas distribution layer facilitates this reaction. One common oxygen catalyst comprises platinum powder very thinly coated onto a carbon paper or cloth. Many designs employ a rough and porous catalyst to increase surface area of the platinum exposed to the hydrogen and oxygen.

In one embodiment, fuel cell 20 comprises a set of bi-polar plates that each includes channel fields on opposite faces that distribute the hydrogen and oxygen. One channel field distributes hydrogen while a channel field on the opposite face distributes oxygen. Multiple bi-polar plates can be stacked to produce a 'fuel cell stack' in which a membrane electrode assembly is disposed between each pair of adjacent bi-polar plates. Since the electrical generation process in fuel cell 20 is exothermic, fuel cell 20 may implement a thermal management system to dissipate heat from the fuel cell. Fuel cell 20 may also employ a number of humidification plates (HP) to manage moisture levels in the fuel cell. Further description of a fuel cell suitable for use with the present invention is included in commonly owned co-pending patent application entitled "Micro Fuel Cell Architecture" naming Ian Kaye as inventor and filed on the same day as this patent application. This application is incorporated by reference for all purposes.

While the present invention will mainly be discussed with respect to PEM fuel cells, it is understood that the present invention may be practiced with other fuel cell architectures. The main difference between fuel cell architectures is the type of ion conductive membrane used. In one embodiment, fuel cell **20** is phosphoric acid fuel cell that employs liquid phosphoric acid for ion exchange. Solid oxide fuel cells employ a hard, non-porous ceramic compound for ion exchange and may be suitable for use with the present invention. Generally, any fuel cell architecture may benefit from fuel processor improvements described herein. Other such fuel cell architectures include direct methanol, alkaline and molten carbonate fuel cells.

Fuel cell 20 generates dc voltage that may be used in a wide variety of applications. For example, electricity generated by
fuel cell 20 may be used to power a motor or light. In one embodiment, the present invention provides 'small' fuel cells that are designed to output less than 200 watts of power (net or total). Fuel cells of this size are commonly referred to as 'micro fuel cells' and are well suited for use with portable
electronics. In one embodiment, fuel cell 20 is configured to generate from about 1 milliwatt to about 200 watts. In another embodiment, fuel cell 20 generates from about 3 W to about 20 W. Fuel cell 20 may also be a stand-alone fuel cell, which is a single unit that produces power as long as it has an a)
oxygen and b) hydrogen or a hydrocarbon fuel supply. A fuel cell 20 that outputs from about 40 W to about 100 W is well suited to power a laptop computer.

FIG. 1B illustrates schematic operation for fuel cell system 10 in accordance with a specific embodiment of the present invention. As shown, fuel cell system 10 comprises fuel container 16, hydrogen fuel source 17, fuel processor 15, fuel cell 20, multiple pumps 21 and fans 35, fuel lines and gas lines, and one or more valves 23. While the present invention will now primarily be described with respect to methanol as fuel source 17, it is understood that the present invention may employ another fuel source 17 such as one provided above.

Fuel container 16 stores methanol as a hydrogen fuel source 17. An outlet 26 of fuel container 16 provides methanol 17 into hydrogen fuel source line 25. As shown, line 25 divides into two lines: a first line 27 that transports methanol 17 to a burner 30 for fuel processor 15 and a second line 29 that transports methanol 17 to reformer 32 in fuel processor **15.** Lines **25, 27** and **29** may comprise plastic tubing, for example. Separate pumps **21***a* and **21***b* are provided for lines **27** and **29**, respectively, to pressurize the lines and transmit the fuel source at independent rates if desired. A model P625 pump as provided by Instech of Plymouth Meeting, Pa. is 5 suitable to transmit liquid methanol for system **10** is suitable in this embodiment. A flow sensor or valve **23** situated on line **29** between storage device **16** and fuel processor **18** detects and communicates the amount of methanol **17** transfer between storage device **16** and suitable control, such as digital control applied by a processor that implements instructions from storage device **16** to reformer **32**.

Fan 35*a* delivers oxygen and air from the ambient room 15 through line 31 to regenerator 36 of fuel processor 15. Fan 35*b* delivers oxygen and air from the ambient room through line 33 to regenerator 36 of fuel processor 15. In this embodiment, a model AD2005DX-K70 fan as provided by Adda USA of California is suitable to transmit oxygen and air for 20 fuel cell system 10. A fan 37 blows cooling air over fuel cell 20 and its heat transfer appendages 46.

Fuel processor 15 receives methanol 17 from storage device 16 and outputs hydrogen. Fuel processor 15 comprises burner 30, reformer 32, boiler 34 and dewar 150. Burner 30 25 includes an inlet that receives methanol 17 from line 27 and a catalyst that generates heat with methanol presence. In one embodiment, burner 30 includes an outlet that exhausts heated gases to a line 41, which transmits the heated gases over heat transfer appendages 46 of fuel cell 20 to pre-heat the 30 fuel cell and expedite warm-up time needed when initially turning on fuel cell 20. An outlet of burner 30 may also exhaust heated gases into the ambient room.

Boiler **34** includes an inlet that receives methanol **17** from line **29**. The structure of boiler **34** permits heat produced in 35 burner **30** to heat methanol **17** in boiler **34** before reformer **32** receives the methanol **17**. Boiler **34** includes an outlet that provides heated methanol **17** to reformer **32**.

Reformer **32** includes an inlet that receives heated methanol **17** from boiler **34**. A catalyst in reformer **32** reacts with the 40 methanol **17** and produces hydrogen and carbon dioxide. This reaction is slightly endothermic and draws heat from burner **30**. A hydrogen outlet of reformer **32** outputs hydrogen to line **39**. In one embodiment, fuel processor **15** also includes a preferential oxidizer that intercepts reformer **32** hydrogen 45 exhaust and decreases the amount of carbon monoxide in the exhaust. The preferential oxidizer and a catalyst based on, for example ruthenium or platinum, that is preferential to carbon monoxide over carbon dioxide. 50

Dewar 150 pre-heats a process gas or liquid before the air enters burner 30. Dewar 150 also reduces heat loss from fuel cell 15 by heating the incoming process liquids or gases before the heat escapes fuel processor 15. In one sense, dewar 150 acts as a regenerator that uses waste heat in fuel processor 55 15 to improve thermal management and thermal efficiency of the fuel processor. Specifically, waste heat from burner 30 may be used to pre-heat incoming air provided to burner 30 to reduce heat transfer to the air in the burner so more heat transfers to reformer 32. Dewar 150 is described in further 60 detail below.

Line **39** transports hydrogen from fuel processor **15** to fuel cell **20**. Gaseous delivery lines **31**, **33** and **39** may comprise polymeric or metallic tubing, for example. A hydrogen flow sensor (not shown) may also be added on line **39** to detect and 65 communicate the amount of hydrogen being delivered to fuel cell **20**. In conjunction with the hydrogen flow sensor and 8

suitable control, such as digital control applied by a processor that implements instructions from stored software, fuel processor **15** regulates hydrogen gas provision to fuel cell **20**.

Fuel cell **20** includes an hydrogen inlet port that receives hydrogen from line **39** and delivers it to a hydrogen intake manifold for delivery to one or more bi-polar plates and their hydrogen distribution channels. An oxygen inlet port of fuel cell **20** receives oxygen from line **33** and delivers it to an oxygen intake manifold for delivery to one or more bi-polar plates and their oxygen distribution channels. An anode exhaust manifold collects gases from the hydrogen distribution channels and delivers them to an anode exhaust port, which outlets the exhaust gases into the ambient room. A cathode exhaust manifold collects gases from the oxygen distribution channels and delivers them to a cathode exhaust port.

The schematic operation for fuel cell system 10 shown in FIG. 1B is exemplary and other variations on fuel cell system design, such as reactant and byproduct plumbing, are contemplated. In addition to the components shown in shown in FIG. 1B, system 10 may also include other elements such as electronic controls, additional pumps and valves, added system sensors, manifolds, heat exchangers and electrical interconnects useful for carrying out functionality of system 10 that are known to one of skill in the art and omitted herein for sake of brevity.

FIG. 1C illustrates an embodiment of fuel system 10 that routes unused hydrogen from fuel cell 20 back to burner 30. Burner 30 includes a catalyst that reacts with the unused hydrogen to produce heat. Since hydrogen consumption within fuel cell 20 is often incomplete and the anode exhaust often includes unused hydrogen, re-routing the anode exhaust to burner 30 allows fuel cell system 10 to capitalize on unused hydrogen in fuel cell 20 and increase hydrogen usage and efficiency in system 10. As the term is used herein, unused hydrogen generally refers to hydrogen output from a fuel cell.

Line 51 is configured to transmit unused hydrogen from fuel cell 20 to burner 30 of fuel processor 15. For FIG. 1C, burner 30 includes two inlets: an inlet 55 configured to receive the hydrogen fuel source 17 and an inlet 53 configured to receive the hydrogen from line 51. Anode gas collection channels, which distribute hydrogen from fuel processor 15 to each membrane electrode assembly layer, collect and exhaust the unused hydrogen. An inlet fan pressurizes line 39 that delivers the hydrogen from an outlet of fuel processor 15 to an anode inlet of fuel cell 20. The inlet fan also pressurizes the anode gas collection channels for distribution of hydrogen within fuel cell 20. In one embodiment, gaseous delivery in line 51 back to fuel processor 15 relies on pressure at the exhaust of the anode gas distribution channels, e.g., in the anode exhaust manifold. In another embodiment, an extra fan is added to line 51 to pressurize line 51 and return unused hydrogen back to fuel processor 15.

Burner 30 also includes an inlet 59 configured to receive oxygen from an oxygen exhaust included in fuel cell 20. Cathode gas collection channels, which distribute oxygen and air from the ambient room to each membrane electrode assembly layer, collect and exhaust the unused oxygen. Line 61 delivers unused oxygen from an exhaust manifold, which collects oxygen from each cathode gas collection channel, to inlet 59. Burner 30 thus includes two oxygen inlets: inlet 59 and an inlet 57 configured to receive oxygen from the ambient room after delivery though dewar 150. Since oxygen consumption within fuel cell 20 is often incomplete and the cathode exhaust includes unused oxygen, re-routing the cathode exhaust to burner 30 allows fuel cell system 10 to capitalize on unused oxygen in fuel cell 20 and increase oxygen usage and efficiency in system 10.

In one embodiment, fuel processor 15 is a steam reformer that only needs steam to produce hydrogen. Several types of reformers suitable for use in fuel cell system 10 include steam reformers, auto thermal reformers (ATR) or catalytic partial oxidizers (CPOX). ATR and CPOX reformers mix air with the fuel and steam mix. ATR and CPOX systems reform fuels such as methanol, diesel, regular unleaded gasoline and other hydrocarbons. In a specific embodiment, storage device 16 provides methanol 17 to fuel processor 15, which reforms the methanol at about 250° C. or less and allows fuel cell system 10 use in applications where temperature is to be minimized.

FIG. 2A illustrates a cross-sectional side view of fuel pro- 15 cessor 15 in accordance with one embodiment of the present invention. FIG. 2B illustrates a cross-sectional front view of fuel processor 15 taken through a mid-plane of processor 15 that also shows features of end plate 82. Fuel processor 15 reforms methanol to produce hydrogen. Fuel processor 15 20 34. Walls 111 in monolithic structure 100 (see cross section in comprises monolithic structure 100, end plates 82 and 84, reformer 32, burner 30, boiler 34, boiler 108, dewar 150 and housing 152. Although the present invention will now be described with respect to methanol consumption for hydrogen production, it is understood that fuel processors of the ²⁵ present invention may consume another fuel source, as one of skill in the art will appreciate.

As the term is used herein, 'monolithic' refers to a single and integrated structure that includes at least portions of $_{30}$ multiple components used in fuel processor 15. As shown, monolithic structure 100 includes reformer 32, burner 30, boiler 34 and boiler 108. Monolithic structure 100 may also include associated plumbing inlets and outlets for reformer 32, burner 30 and boiler 34. Monolithic structure 100 com-35 prises a common material 141 that constitutes the structure. Common material 141 is included in walls that define the reformer 32, burner 32 and boilers 34 and 108. Specifically, walls 111, 119, 120, 122, 130, 132, 134 and 136 all comprise common material 141. Common material 141 may comprise 40 a metal, such as copper, silicon, stainless steel, inconel and other metal/alloys displaying favorable thermal conducting properties. The monolithic structure 100 and common material 141 simplify manufacture of fuel processor 15. For example, using a metal for common material 141 allows 45 monolithic structure 100 to be formed by extrusion or casting. In some cases, monolithic structure 100 is consistent in cross sectional dimensions between end plates 82 and 84 and solely comprises copper formed in a single extrusion. Common material 141 may also include a ceramic, for example. A ceramic monolithic structure 100 may be formed by sintering.

Housing 152 provides mechanical protection for internal components of fuel processor 15 such as burner 30 and reformer 32. Housing 152 also provides separation from the environment external to processor 15 and includes inlet and 55 outlet ports for gaseous and liquid communication in and out of fuel processor 15. Housing 152 includes a set of housing walls 161 that at least partially contain a dewar 150 and provide external mechanical protection for components in fuel processor 15. Walls 161 may comprises a suitably stiff $_{60}$ material such as a metal or a rigid polymer, for example. Dewar 150 improves thermal heat management for fuel processor 15 and will be discussed in further detail with respect to FIG. 4A.

Together, monolithic structure 100 and end plates 82 and 65 84 structurally define reformer 32, burner 30, boiler 34 and boiler 108 and their respective chambers. Monolithic struc-

ture 100 and end plates 82 and 84 are shown separate in FIG. 2A for illustrative purposes, while FIG. 4A shows them together.

Referring to FIG. 2B, boiler 34 heats methanol before reformer 32 receives the methanol. Boiler 34 receives the methanol via fuel source inlet 81, which couples to the methanol supply line 27 of FIG. 1B. Since methanol reforming and hydrogen production via a catalyst 102 in reformer 32 often requires elevated methanol temperatures, fuel processor 15 pre-heats the methanol before receipt by reformer 32 via boiler 34. Boiler 34 is disposed in proximity to burner 30 to receive heat generated in burner 30. The heat transfers via conduction through monolithic structure from burner 30 to boiler 34 and via convection from boiler 34 walls to the methanol passing therethrough. In one embodiment, boiler 34 is configured to vaporize liquid methanol. Boiler 34 then passes the gaseous methanol to reformer 32 for gaseous interaction with catalyst 102.

Reformer 32 is configured to receive methanol from boiler FIG. 3A) and end walls 113 (FIG. 2B) on end plates 82 and 84 define dimensions for a reformer chamber 103. In one embodiment, end plate 82 and/or end plate 84 includes also channels 95 (FIG. 2A) that route heated methanol exhausted from boiler 34 into reformer 32. The heated methanol then enters the reformer chamber 103 at one end of monolithic structure 100 and passes to the other end where the reformer exhaust is disposed. In another embodiment, a hole disposed in a reformer 32 wall receives inlet heated methanol from a line or other supply. The inlet hole or port may be disposed on a suitable wall 111 or 113 of reformer 32.

Reformer 32 includes a catalyst 102 that facilitates the production of hydrogen. Catalyst 102 reacts with methanol 17 and facilitates the production of hydrogen gas and carbon dioxide. In one embodiment, catalyst 102 comprises pellets packed to form a porous bed or otherwise suitably filled into the volume of reformer chamber 103. In one embodiment, pellet sizes are designed to maximize the amount of surface area exposure to the incoming methanol. Pellet diameters ranging from about 50 microns to about 1.5 millimeters are suitable for many applications. Pellet diameters ranging from about 300 microns to about 1500 microns are suitable for use with reformer chamber 103. Pellet sizes and packing may also be varied to control the pressure drop that occurs through reformer chamber 103. In one embodiment, pressure drops from about 0.2 to about 5 psi gauge are suitable between the inlet and outlet of reformer chamber 103. Pellet sizes may be varied relative to the cross sectional size of reformer chamber 103, e.g., as reformer chamber 103 increases in size so may catalyst 102 pellet diameters. In one embodiment, the ratio of pellet diameter (d) to cross sectional height 117 (D) may range from about 0.0125 to about 1. A D/d ratio from about 5 to about 20 is also suitable for many applications. A packing density may also characterize packing of catalyst 102 in reformer chamber 103. For a copper zinc catalyst 102, packing densities from about 0.3 grams/milliliter to about 2 grams/ milliliter are suitable. Packing densities from about 0.9 grams/milliliter to about 1.4 grams/milliliter are appropriate for the embodiment shown in FIG. 3A.

One suitable catalyst 102 may include CuZn on alumina pellets when methanol is used as a hydrocarbon fuel source 17. Other materials suitable for catalyst 102 may be based on nickel, platinum, palladium, or other precious metal catalysts either alone or in combination, for example. Catalyst 102 pellets are commercially available from a number of vendors known to those of skill in the art. Pellet catalysts may also be disposed within a baffling system disposed in the reformer chamber **103**. The baffling system includes a set of walls that guide the fuel source along a non-linear path. The baffling slows and controls flow of gaseous methanol in chamber **103** to improve interaction between the gaseous methanol and pellet catalyst **102**. Catalyst **102** may alternatively comprise 5 catalyst materials listed above coated onto a metal sponge or metal foam. A wash coat of the desired metal catalyst material onto the walls of reformer chamber **103** may also be used for reformer **32**.

Reformer **32** is configured to output hydrogen and includes 10 an outlet port **87** that communicates hydrogen formed in reformer **32** outside of fuel processor **15**. In fuel cell system **10**, port **87** communicates hydrogen to line **39** for provision to hydrogen distribution **43** in fuel cell **20**. Port **87** is disposed on a wall of end plate **82** and includes a hole that passes through 15 the wall (see FIG. 2B). The outlet hole port may be disposed on any suitable wall **111** or **113**.

Hydrogen production in reformer 32 is slightly endothermic and draws heat from burner 30. Burner 30 generates heat and is configured to provide heat to reformer 32. Burner 30 is 20 disposed annularly about reformer 32, as will be discussed in further detail below. As shown in FIG. 2B, burner 30 comprises two burners (or burner sections) 30a and 30b and their respective burner chambers 105a and 105b that surround reformer 32. Burner 30 includes an inlet that receives metha- 25 nol 17 from boiler 108 via a channel in one of end plates 82 or 84. In one embodiment, the burner inlet opens into burner chamber 105a. The methanol then travels the length 142 of burner chamber 105a to channels disposed in end plate 82 that route methanol from burner chamber 105a to burner chamber 30 **105***b*. The methanol then travels the back through the length 142 of burner chamber 105b to burner exhaust 89. In another embodiment, the burner inlet opens into both chambers 105a and 105b. The methanol then travels the length 142 of both chambers 105a and 105b to burner exhaust 89.

In one embodiment, burner 30 employs catalytic combustion to produce heat. A catalyst 104 disposed in each burner chamber 105 helps a burner fuel passed through the chamber generate heat. In one embodiment, methanol produces heat in burner 30 and catalyst 104 facilitates the methanol production 40 of heat. In another embodiment, waste hydrogen from fuel cell 20 produces heat in the presence of catalyst 104. Suitable burner catalysts 104 may include platinum or palladium coated onto a suitable support or alumina pellets for example. Other materials suitable for catalyst 104 include iron, tin 45 oxide, other noble-metal catalysts, reducible oxides, and mixtures thereof. The catalyst 104 is commercially available from a number of vendors known to those of skill in the art as small pellets. The pellets that may be packed into burner chamber 105 to form a porous bed or otherwise suitably filled into the 50 burner chamber volume. Catalyst 104 pellet sizes may be varied relative to the cross sectional size of burner chamber 105. Catalyst 104 may also comprise catalyst materials listed above coated onto a metal sponge or metal foam or wash coated onto the walls of burner chamber 105. A burner outlet 55 port 89 (FIG. 2A) communicates exhaust formed in burner 30 outside of fuel processor 15.

Some fuel sources generate additional heat in burner **30**, or generate heat more efficiently, with elevated temperatures. Fuel processor **15** includes a boiler **108** that heats methanol ⁶⁰ before burner **30** receives the fuel source. In this case, boiler **108** receives the methanol via fuel source inlet **85**. Boiler **108** is disposed in proximity to burner **30** to receive heat generated in burner **30**. The heat transfers via conduction through monolithic structure from burner **30** to boiler **108** and via convection from boiler **108** walls to the methanol passing therethrough.

Air including oxygen enters fuel processor 15 via air inlet port 91. Burner 30 uses the oxygen for catalytic combustion of methanol. As will be discussed in further detail below with respect to FIGS. 4A and 4B, air first passes along the outside of dewar 150 before passing through apertures in the dewar and along the inside of dewar 150. This heats the air before receipt by air inlet port 93 of burner 30.

FIG. 3A illustrates a cross-sectional front view of monolithic structure 100 as taken through a mid-plane 121 in accordance with one embodiment of the present invention. Monolithic structure 100 extends from end plate 82 to end plate 84. The cross section of monolithic structure 100 shown in FIG. 3A extends from one end of structure 100 at end plate 82 to the other end of structure 100 at end plate 84. Monolithic structure 100 includes reformer 32, burner 30, boiler 34 and boiler 108 between end plates 82 and 84.

Reformer 32 includes a reformer chamber 103, which is a voluminous space in fuel processor 15 that includes the reforming catalyst 102, opens to the fuel source inlet (from boiler 34 for fuel processor 15), and opens to hydrogen outlet 87. Side walls 111 define a non-planar cross-sectional shape for reformer 32 and its reformer chamber 103. Walls 113 on end plates 82 and 84 close the reformer chamber 103 on either end of the chamber 103 and include the inlet and outlet ports to the chamber 103.

Reformer chamber 103 includes a non-planar volume. As the term is used herein, a non-planar reformer chamber 103 refers to a shape in cross section that is substantially non-flat or non-linear. A cross section refers to a planar slice that cuts 30 through the fuel processor or component. For cross sections that include multiple fuel processor components (e.g., both burner 30 and reformer 32), the cross section includes both components. For the vertical and front cross section 121 shown in FIG. 3A, the cross section dimensions shown are 35 consistent for monolithic structure 100 from end plate 82 to end plate 84, and are consistent at each cross section 121 (FIG. 2A).

Reformer 32 and its reformer chamber 103 may employ a quadrilateral or non-quadrilateral cross-sectional shape. Four sides define a quadrilateral reformer chamber 103 in cross section. Four substantially orthogonal sides define rectangular and square quadrilateral reformers 32. A non-quadrilateral reformer 32 may employ cross-sectional geometries with more or less sides, an elliptical shape (see FIG. 3B), and more complex cross-sectional shapes. As shown in FIG. 3A, reformer 32 includes a six-sided cross-sectional 'P-shape' with chamfered corners. One corner section of reformer 32 is removed from monolithic structure 100 to allow for boiler 34 proximity to burner 30.

Reformer chamber 103 is characterized by a cross-sectional width 115 and a cross-sectional height 117. A maximum linear distance between inner walls 111 of chamber 103 in a direction spanning a cross section of reformer chamber 103 quantifies cross-sectional width 155. A maximum linear distance between inner walls 111 of chamber 103 orthogonal to the width 115 quantifies cross-sectional height 117. As shown, cross-sectional height 117 is greater than one-third the cross-sectional width 115. This height/width relationship increases the volume of reformer chamber 103 for a given fuel processor 15. In one embodiment, cross-sectional height 117 is greater than one-half cross-sectional width 115. In another embodiment, cross-sectional height 117 is greater than the cross-sectional width 115.

Referring back to FIG. 2A, reformer chamber 103 includes a length 142 (orthogonal to the width 115 and height 117) that extends from one end of monolithic structure 100 at end plate 82 to the other end of structure 100 at end plate 84. In one embodiment, reformer chamber 103 has a length 142 to width 115 ratio less than 20:1. In a less elongated design, reformer chamber 103 has a length 142 to width 115 ratio less than 10:1.

Reformer 32 provides a voluminous reformer chamber 5 103. This three dimensional configuration for reformer chamber 103 contrasts micro fuel processor designs where the reformer chamber is etched as micro channels onto a planar substrate. The non-planar dimensions of reformer chamber 103 permit greater volumes for reformer 32 and permit more 10 catalyst 102 for a given size of fuel processor 15. This increases the amount of methanol that can be processed and increases hydrogen output for a particular fuel processor 15 size. Reformer 32 thus improves fuel processor's 15 suitability and performance in portable applications where fuel pro- 15 cessor size is important or limited. In other words, since the size of inlet and outlet plumbing and ports varies little while increasing the reformer chamber 103 volume, this allows fuel processor 15 to increase hydrogen output and increase power density for portable applications while maintaining size and 20 weight of the associated plumbing relatively constant. In one embodiment, reformer chamber 103 comprises a volume greater than about 0.1 cubic centimeters and less than about 50 cubic centimeters. In some embodiments, reformer 32 volumes between about 0.5 cubic centimeters and about 2.5 25 cubic centimeters are suitable for laptop computer applications.

Fuel processor 15 includes at least one burner 30. Each burner 30 includes a burner chamber 105. For a catalytic burner 30, the burner chamber 105 is a voluminous space in 30 fuel processor 15 that includes catalyst 104. For communication or burner reactants and products to and from the burner chamber 105, the burner chamber 105 may directly or indirectly open to a fuel source inlet (from boiler 108 for fuel processor 15), open to an air inlet 93, and open to a burner 35 exhaust 89.

The number of burners 30 and burner chambers 105 may vary with design. Monolithic structure 100 of FIG. 3A includes a dual burner 30a and 30b design having two burner chambers 105a and 105b, respectively, forming non-continu- 40 ous chambers that substantially surround reformer 32 in cross section. Burner 30a comprises side walls 119a (FIG. 3A) included in monolithic structure 100 and end walls 113 on end plates 82 and 84 (FIG. 2B) that define burner chamber 105a. Similarly, burner 30b includes side walls 119b (FIG. 3A) 45 included in monolithic structure 100 and end walls 113 on end plates 82 and 84 (FIG. 2B) that define burner chamber 105b. Monolithic structure 100 of FIG. 3C includes a single burner 30c with a single burner chamber 105c that fully surrounds reformer 32. Tubular arrangement of FIG. 3B includes over 50 forty burners 204 that fully surround reformer 202. Monolithic structure 452 of FIG. 4F includes a single burner divided into 104 burner chambers that fully surround reformer 32.

Referring to FIG. 2B, each burner 30 is configured relative 55 to reformer 32 such that heat generated in a burner 30 transfers to reformer 32. In one embodiment, the one or more burners 30 are annularly disposed about reformer 32. As the term is used herein, annular configuration of at least one burner 30 relative to reformer 32 refers to the burner 30 60 having, made up of, or formed by, continuous or non-continuous segments or chambers 105 that surround reformer 32. The annular relationship is apparent in cross section. For burner and reformer arrangements, surrounding refers to a burner 30 bordering or neighboring the perimeter of reformer 32 such 65 that heat may travel from a burner 30 to the reformer 32. Burners 30*a* and 30*b* may surround reformer 32 about the

perimeter of reformer 32 to varying degrees based on design. At the least, one or more burners 30 surround greater than 50 percent of the reformer 32 cross-sectional perimeter. This differentiates fuel processor 15 from planar and plate designs where the burner and reformer are co-planar and of similar dimensions, and by geometric logic, the burner neighbors less than 50 percent of the reformer perimeter. In one embodiment, one or more burners 30 surround greater than 75 percent of the reformer 32 cross-sectional perimeter. Increasing the extent to which burner 30 surrounds reformer 32 perimeter in cross section increases the surface area of reformer 32 that can be used to heat the reformer volume via heat generated in the burner. For some fuel processor 15 designs, one or more burners 30 may surround greater than 90 percent of the reformer 32 cross-sectional perimeter. For the embodiment shown in FIG. 3B, burner 30 surrounds the entire reformer 32 cross-sectional perimeter.

Although the present invention will now be described with respect to burner 30 annularly disposed about reformer 32, it is understood that monolithic structure 100 may comprise the reverse configuration. That is, reformer 32 may be annularly disposed about burner 30. In this case, reformer 32 may comprise one or more continuous or non-continuous segments or chambers 103 that surround burner 30.

In one embodiment, each burner **30** and its burner chamber **105** has a non-planar cross-sectional shape. A non-planar burner **30** may employ cross-sectional shapes such as quadrilaterals, non-quadrilateral geometries with more or less sides, an elliptical shape (see FIG. **3**B for circular/tubular burners **30**), or more complex cross-sectional shapes. As shown in FIG. **3**A, each burner **30** includes a six-sided crosssectional 'L' shape (with chamfered corners) that bends 90 degrees about reformer **32**.

Each burner 30 thus bilaterally borders reformer 32. N-lateral bordering in this sense refers to the number of sides, N, of reformer 32 that a burner 30 (and its burner chamber 105) borders in cross section. Thus, burner 30b borders the right and bottom sides of reformer 32, while burner 30a borders the top and left sides of reformer 32. A 'U-shaped' burner 30 may be employed to trilaterally border reformer 32 on three sides. Together, burners 30a and 30b quadrilaterally border reformer 32 on all four orthogonal reformer 32 sides. The reformer 32 used in the configuration of FIG. 3B includes multiple tubular burners that quadrilaterally border reformer 32. FIG. 3C illustrates a cross-sectional front view of monolithic structure 100 that comprises a single burner 30c having an 'O-shape' that completely surrounds reformer chamber 103 in accordance with one embodiment of the present invention. Burner 30c is a continuous chamber about the perimeter of reformer 32 and quadrilaterally borders reformer 32.

Heat generated in burner 30 transfers directly and/or indirectly to reformer 32. For the monolithic structure 100 of FIG. 3A, each burner 30 and reformer 32 share common walls 120 and 122 and heat generated in each burner 30 transfers directly to reformer 32 via conductive heat transfer through common walls 120 and 122. Wall 120 forms a boundary wall for burner 30b and a boundary wall for reformer 32. As shown, one side of wall 120 opens to burner chamber 105bwhile another portion of the wall opens to reformer chamber 103. Wall 120 thus permits direct conductive heat transfer between burner 30b and reformer 32. Similarly, wall 122 forms a boundary wall for burner 30a and a boundary wall for reformer 32, opens to burner chamber 105a, opens to reformer chamber 103, and permits direct conductive heat transfer between burner 30a and reformer 32. Walls 120 and 122 are both non-planar in cross section and border multiple sides of reformer chamber 103 that are neighbored by burners 30b and 30a. Wall 120 thus provides direct conductive heat transfer in multiple orthogonal directions 128 and 129 from burner 30a to reformer 32. Wall 122 similarly provides direct conductive heat transfer in directions opposite to 128 and 129 from burner 30*b* to reformer 32.

Boiler 34 comprises cylindrical walls 143 included in monolithic structure 100 and end walls 113 on end plates 82 and 84 (see FIG. 2B) that define boiler chamber 147. Circular walls 143 in cross section form a cylindrical shape for boiler 34 that extends from routing end 82 to routing end 84. Boiler 34 is disposed in proximity to burners 30a and 30b to receive heat generated in each burner 30. For monolithic structure 100, boiler 34 shares a common wall 130 with burner 30a and a common wall 132 with burner 30b. Common walls 130 and 132 permit direct conductive heat transfer from each burner 15 30 to boiler 34. Boiler 34 is also disposed between burners 30 and reformer 32 to intercept thermal conduction consistently moving from the high temperature and heat generating burners 30 to the endothermic reformer 32.

Boiler 108 is configured to receive heat from burner 30 to 20 heat methanol before burner 30 receives the methanol. Boiler 108 also comprises a tubular shape having a circular cross section that extends through monolithic structure 100 from end plate 82 to end plate 84. Boiler 108 is disposed in proximity to burners 30a and 30b to receive heat generated in each 25 burner 30, which is used to heat the methanol. Boiler 108 shares a common wall 134 with burner 30a and a common wall 136 with burner 30b. Common walls 134 and 136 permit direct conductive heat transfer from burners 30a and 30b to boiler 108.

FIG. 3D illustrates an outside view of end plate 82 in accordance with one embodiment of the present invention. End plate 82 includes fuel source inlet 81, fuel source fuel source inlet 85, hydrogen outlet port 87 and burner air inlet 93. Fuel source inlet 81 includes a hole or port in end wall 113 35 of end plate 82 that communicates methanol (usually as a liquid) from an external methanol supply to boiler 34 for heating the methanol before receipt by reformer 32. Methanol fuel source inlet 85 includes a hole or port in end wall 113 of end plate 82 that communicates methanol (usually as a liquid) 40 from an external methanol supply to boiler 108 for heating the methanol before receipt by burner 30. Burner air inlet 93 includes a hole or port in end wall 113 of end plate 82 that communicates air and oxygen from the ambient room after it has been preheated in dewar 150. Hydrogen outlet port 87 45 communicates gaseous hydrogen from reforming chamber 103 outside fuel processor 15.

Bolt holes 153 are disposed in wings 145 of monolithic structure 100. Bolt holes 153 permit the passage of bolts therethrough and allow securing of structure 100 and end 50 plates 82 and 84.

FIG. 3B illustrates a cross-sectional layout of a tubular design 200 for use in fuel processor 15 in place of monolithic structure 100 in accordance with another embodiment of the present invention. Structure 200 includes a reformer 202, 55 burner 204, boiler 206 and boiler 208.

The cross-sectional design 200 shown in FIG. 3B is consistent throughout a cylindrical length between end plates (not shown) that include inlet and outlet ports for supply and exhaust of gases to components of design 200. The circular 60 shape of reformer 202, burner chambers 212, boiler 206 and boiler 208 thus extends for the entire cylindrical length between the end plates. The end plates may also be responsible for routing gases between individual tubes, such as between tubular burners 234.

Reformer 202 includes cylindrical walls 203 that define a substantially circular cross section. Reformer 232 thus resembles a hollow cylinder in three dimensions that defines a tubular reformer chamber 210. In general, reformer 202 may include any elliptical shape (a circle represents an ellipse of about equal orthogonal dimensions) suitable for containing the catalyst 102, for methanol flow through reformer chamber 210, for hydrogen production in reformer chamber 210, and for hydrogen flow in reformer chamber 210. As shown, reformer chamber 210 is defined by a cross sectional width and a cross sectional height that are substantially equal and thus reformer 202 includes a 1:1 cross sectional aspect ratio.

Burner 204 comprises a set of cylindrical walls 214 that each defines a tubular burner chamber 212. As shown, tubular design 200 includes over forty tubular burner chambers 212 that fully surround the cross sectional perimeter of reformer 32. Each tubular burner chamber 212 includes a substantially circular cross-section defined by the cylindrical wall 214. Each tubular burner chamber 212 includes catalyst 104 that facilitates heat generation from methanol. Burner 204 may comprise from about two to about two hundred cylindrical walls 214 and tubular burner chambers 212. Some designs may include from about ten to about sixty tubular burner chambers 212. In one embodiment, each cylindrical wall 214 comprises a metal and is extruded to its desired dimensions. In a specific embodiment, cylindrical wall 214 comprises nickel. The nickel wall 214 may be formed by electroplating nickel onto a suitable substrate such as zinc or aluminum that may subsequently etched out to leave the nickel tube. Other materials that a nickel wall 214 may be formed onto include zinc, tin, lead, wax or plastics. In addition to nickel, wall 214 may include gold, silver, copper, stainless steel, ceramics and materials that display suitable thermal properties without causing complications with burner catalyst 104.

As shown, burner 204 fully annularly surrounds the crosssectional perimeter of reformer 202. In this case, burner 204 comprises three ring-like layers 216, 218 and 220 of tubes 214 disposed circularly about reformer 202 and at three different radii. The tubes 214 in each layer 216, 218 and 220 circumscribe reformer 202. Heat generated in each tubular chamber 212 of burner 30 transfers directly or indirectly to reformer 202 via several paths: a) heat conduction through the tubes 214 in layer 216 to the walls of reformer 202; b) heat conduction through the tubes 214 in outer layers 218 and 220 to tubes 214 in layer 216 and to the walls of reformer 202; and/or c) heat radiation between tubes 214 in outer layers 218 and 220 and tubes 214 and then conduction inward to reformer 202.

Boiler 206 is configured to heat methanol before reformer 202 receives the methanol. Boiler 206 receives heat from burner 204 and comprises a cylindrical wall 207 that defines a tubular shape for the boiler. Boiler 206 is disposed in proximity to burner tubes 214 to receive heat generated in each burner chamber 212. Specifically, boiler 206 is disposed in the second ring-like layer 218 and receives heat from adjacent burner chambers 212 in layers 216, 218 and 220. Burner 204 provides heat to boiler 206 via conduction through the walls of each adjacent tube 214 and through wall 207.

Boiler 208 is configured to heat methanol before burner 204 receives the methanol. Boiler 208 receives heat from burner 204 and also comprises a cylindrical wall 209 that defines a tubular shape for the boiler. Similar to boiler 206, boiler 208 is disposed in the second ring-like layer 218 and receives heat from adjacent burner chambers 212 in layers 216. 218 and 220.

In one embodiment, a monolithic fuel processor 15 comprises multiple segments joined together in the direction of gas flow in reformer chamber 103 and joined at sectional lines 121. Each segment has a common profile as shown in FIG. 3A

65

and may comprise metal or ceramic elements that are bonded or brazed perpendicular to the direction of gas flow. Alternatively, fuel processor 15 may comprise a single long monolithic piece that bounds all of reformer 32, burner 30, boiler 34 and boiler 108 except for areas bound by end pieces 82 and 5 84.

In another embodiment, fuel processor **15** comprises multiple pieces joined together in cross section. FIG. 3E illustrates a fuel processor **15** in accordance with another embodiment of the present invention. In this case, fuel processor **15** 10 comprises three pieces: lower piece **280**, middle piece **282** and cap piece **284**. Lower piece **280** and middle piece **282** attach to form reformer **32** and two burner chambers **30**. Cap piece **284** and middle piece **282** attach to form boilers **34** and **108**. Each piece **208**, **282** and **284** comprises a common 15 material and may be extruded or cast to suitable dimensions. Attachment between the pieces may comprise chemical bonding, for example.

FIG. 4A illustrates a side cross-sectional view of fuel processor 15 and movement of air created by dewar 150 in 20 accordance with one embodiment of the present invention. FIG. 4B illustrates a front cross-sectional view of fuel processor 15 and demonstrates thermal management benefits gained by dewar 150. While thermal management techniques described herein will now be described as fuel processor 25 components, those skilled in the art will recognize that the present invention encompasses methods of thermal management for general application.

A burner **30** in fuel processor **15** generates heat and typically operates at an elevated temperature. Burner **30** operating 30 temperatures greater than 200 degrees Celsius are common. Standards for the manufacture of electronics devices typically dictate a maximum surface temperature for a device. Electronics devices such as laptop computers often include cooling, such as a fan or cooling pipe, to manage and dissipate 35 internal heat. A fuel processor internal to an electronics device that loses heat into the device calls upon the device's cooling system to handle the lost heat.

In one embodiment, fuel processor 15 comprises a dewar 150 to improve thermal management for fuel processor 15. 40 Dewar 150 at least partially thermally isolates components internal to housing 152—such as burner 30—and contains heat within fuel processor 15. Dewar 150 reduces heat loss from fuel processor 15 and helps manage the temperature gradient between burner 30 and outer surface of housing 152. 45 And as will be described below, dewar 150 also pre-heats air before it is received by burner 30.

Dewar 150 at least partially contains burner 30 and reformer 32, and includes a set of dewar walls 154 that help form a dewar chamber 156 and a chamber 158. In some 50 embodiments, dewar 150 fully surrounds burner 30 and reformer 32 in a cross sectional view and at both ends of burner 30 and reformer 32. Less containment by dewar 150 is also suitable to provide thermal benefits described herein. The multipass dewar 300 of FIG. 4E only partially encloses 55 burner 30 and reformer 32 in cross section. In some cases, dewar 150 does not extends fully along the length of mono-lithic structure 100 and provides less than full containment.

As shown in FIG. 4B, dewar 150 annularly surrounds burner 30 in cross section. The set of walls 154 includes side 60 walls 154*a* and 154*c* that combine with top and bottom walls 154*b* and 154*d* to form the rectangular cross section shown in FIG. 4B; and includes two end walls 154*e* and 154*f* that combine with top and bottom walls 154*b* and 154*d* to form the rectangular cross section shown in FIG. 4A. End wall 154*f* 65 includes apertures that permit the passage of inlet and outlet ports 85, 87 and 89 therethough.

Dewar chamber **156** is formed within dewar walls **154** and comprises all space within the dewar walls **154** not occupied by monolithic structure **100**. As shown in FIG. **4B**, dewar chamber **156** surrounds monolithic structure **100**. As shown in FIG. **4B**, chamber **156** comprises ducts between monolithic structure **100** and walls **154** on all four sides of dewar **150**. In addition, chamber **156** comprises air pockets between end walls of dewar **150** and outside surfaces of end plates **82** and **84** on both ends of monolithic structure **100** (FIG. **4**A).

Chamber 158 is formed outside dewar walls 154 between dewar 150 and housing 152. Chamber 158 comprises all space within housing 152 not occupied by dewar 150. As shown in FIG. 4B, housing 152 encloses dewar 150 and the further internal monolithic structure 100. Chamber 158 comprises ducts between walls 154 on all four sides of dewar 150 and housing 152. In addition, chamber 158 comprises air pockets 167 between dewar 150 and housing 152 on both ends that prevent contact and conductive heat transfer between dewar 150 and housing 152 (FIG. 4A).

Dewar 150 is configured such that a process gas or liquid passing through dewar chamber 156 receives heat generated in burner 30. The process gas or liquid may include any reactant used in fuel processor such as oxygen, air, or fuel source 17, for example. Dewar 150 offers thus two functions for fuel processor 15: a) it permits active cooling of components within fuel processor 15 before the heat reaches an outer portion of the fuel processor, and b) it pre-heats the air going to burner 30. For the former, air moves through fuel processor 15 and across walls 154 of dewar 150 such that the cooler air absorbs heat from the warmer fuel processor 15 components.

As shown in FIG. 4A, housing 152 includes an air inlet port 91 or hole that permits the passage of air from outside housing 152 into air into chamber 158. A fan usually provides the air directly to fuel processor 15 and pressurizes the air coming through port 91. Top and bottom walls 154b and 154d include air inlet ports or holes 172 that allow air to pass from chamber 158 to dewar chamber 156. Air flow through fuel processor 15 then flows: in air inlet port 91, through chamber 158 along the length of the dewar 150, through holes 172 in walls 154b and 154d, through chamber 156 back along the length of the dewar 150 in the opposite direction as in through chamber 158, and into air inlet ports 176 that allow the air to enter burner 30. In chamber 158, the air a) moves across the outside surface of dewar walls 154 and absorbs heat convectively from dewar walls 154, and b) moves across the inside surface of housing 152 and absorbs heat convectively from the housing 152 walls (when housing 152 is at a greater temperature than the air). In chamber 156, the air a) moves across the outside surface of monolithic structure 100 and absorbs heat convectively from the walls of monolithic structure 100, and b) moves across the inside surface of dewar 150 and absorbs heat convectively from dewar walls 154.

Dewar 150 is thus configured such that air passing through the dewar receives heat generated in burner 30 via direct convective heat transfer from walls in monolithic structure 100 on the outside of burner 30 to air passing through dewar chamber 156. Dewar 150 is also configured to such that air passing through chamber 156 receives heat indirectly from burner 30. Indirectly in this sense refers to heat generated in burner 30 moving to another structure in fuel processor 15 before receipt by the air.

FIG. 4C illustrates a thermal diagram of the heat path produced by a wall 154 of dewar 150. Heat from burner 30 conducts through monolithic structure 100 to a surface of structure 100 that opens into dewar chamber 156. From here, the heat a) conducts into the air passing through dewar chamber 156, thereby heating the air; b) radiates to the inner wall

20

25

35

45

155 of dewar wall 154, from which the heat convects into the air passing through dewar chamber 156; c) radiates to the inner wall 155 of dewar wall 154, conducts through wall 154 to the outer surface 157 of dewar wall 154, from which the heat convects into the air passing through dewar chamber 158, 5 and d) radiates to the inner wall 155 of dewar wall 154, conducts through wall 154 to the outer surface 157 of dewar wall 155. So dewar wall 154, conducts through wall 154 to the outer surface 157 of dewar wall 154, conducts through wall 154 to the outer surface 157 of dewar wall 154, conducts through wall 154 to the outer surface 157 of dewar wall 154, radiates to a wall of housing 152, from which the heat convects into the air passing through dewar chamber 158.

Dewar 150 thus provides two streams of convective heat dissipation and active air-cooling in volumes 156 and 158 that prevent heat generated in burner 30 (or other internal parts of fuel processor 15) from escaping the fuel processor.

Reflectance of heat back into chamber 156 decreases the amount of heat lost from fuel processor 150 and increases the heating of air passing through chamber 156. To further improve the radiative reflectance back into chamber 156, an inside surface of dewar wall 154 may include a radiative layer 160 to decrease radiative heat transfer into wall 154 (see FIG. 4B or 4C). Radiative layer 160 is disposed on an inner surface 155 on one or more of walls 154 to increase radiative heat reflectance of the inner surface 155. Generally, the material used in radiative layer 160 has a lower emissivity than the material used in walls 154. Materials suitable for use with walls 154 of dewar 150 include nickel or a ceramic, for example. Radiative layer 160 may comprise gold, platinum, silver, palladium, nickel and the metal may be sputter coated onto the inner surface 155. Radiative layer 160 may also include a low heat conductance. In this case, radiative layer 160 may comprise a ceramic, for example.

When dewar 150 fully encapsulates monolithic structure 100, the dewar then bounds heat loss from the structure and decreases the amount of heat passing out of dewar 150 and housing 152. Fuel processors 15 such as that shown in FIGS. 4A and 4B are well suited to contain heat within housing 152 and manage heat transfer from the fuel processor. In one embodiment, burner operates at a temperature greater than about 200 degrees Celsius and the outer side of the housing remains less than about 50 degrees Celsius. In embodiments for portable applications where fuel processor 15 occupies a small volume, volumes 156 and 158 are relatively small and comprise narrow channels and ducts. In some cases, the height of channels in volumes 156 and 158 is less than 5 millimeters and a wall of burner 30 on monolithic structure is no greater than 10 millimeters from a wall of housing 152.

The thermal benefits gained by use of dewar **150** also permit the use of higher temperature burning fuels as a fuel source for hydrogen production, such as ethanol and gasoline. In one embodiment, the thermal management benefits gained by use of dewar **150** permit reformer **32** to process methanol at temperatures well above 100 degrees Celsius and at temperatures high enough that carbon monoxide production in reformer **32** drops to an amount such that a preferential oxidizer is not needed.

As mentioned above, dewar **150** offers a second function for fuel processor **15** by pre-heating the air going to a burner. Burner **30** relies on catalytic combustion to produce heat. Oxygen in the air provided to burner **30** is consumed as part of the combustion process. Heat generated in the burner **30** will 60 heat cool incoming air, depending on the temperature of the air. This heat loss to incoming cool air reduces the heating efficiency of burner **30**, and typically results in a greater consumption of methanol. To increase the heating efficiency of burner **30**, the present invention heats the incoming air so less heat generated in the burner passes into the incoming air. In other words, chambers and air flow formed by dewar **150**

allow waste heat from the burner to pre-heat air before reaching the burner, thus acting as a regenerator for fuel cell **15**.

While fuel processor 15 of FIGS. 4A and 4B shows dewar 150 encapsulating monolithic structure 100, the present invention may also employ other architectures for dewar 150 and relationships between burner 30 or reformer 32 and dewar 150 that carry out one or both of the dewar functions described above. FIG. 4D illustrates a cross sectional view of a fuel processor 15 that elongates the convective path for cool air flow over a warmer dewar wall 254 in accordance with another embodiment of the present invention. Fuel processor 15 includes a tubular design for the burner 30 and reformer 32.

Dewar 250 routes cool incoming air across an elongated heat transfer path. Dewar 250 includes a spiral wall 254 in cross section that surrounds burner 30 and reformer 32. Spiral wall 254 defines a spiral dewar chamber 256. Cool air enters dewar chamber 256 at a dewar entrance 252. The innermost portion 257 of wall 254 attaches to an outer wall 258 of burner 30. Heat from burner 30 conducts linearly through spiral wall 254. Thus, inner portion 257 is the warmest portion of wall 254, while wall 254 at entrance 252 is typically the coolest. Air progressively warms as it travels through dewar chamber 256. As the air travels inward, temperature of wall 254 rises, as does the amount of heat available for transfer to the air. Depending on the transient temperature of the air, the amount of heat lost from wall 154 may also increase as the air progresses inward.

Spiral dewar **250** elongates convection interaction between the incoming cool air and a wall warmed by the burner. Dewar **250** also increases the number of walls and convective layers in a given radial direction from the fuel processor center. As shown in FIG. **4D**, dewar **250** comprises 4-5 walls and convective layers in a given radial direction, depending on where the number is counted. The number of walls and convective in a radial direction may vary with design. In one embodiment, spiral dewar **250** is configured with from 1 layer to about 50 walls and convective layers in a given radial direction from the fuel processor center. Three layers to 20 layers are suitable for many applications. A channel width **260** defines the duct space between adjacent walls **254**. In one embodiment, channel width **260** ranges from about ¹/₄ millimeters to about 5 millimeters.

Spiral dewar **250** may be constructed by electroplating nickel onto a removable layer such as aluminum or zinc. FIG. **4**H illustrates spiral dewar **250** in an unrolled form during initial construction in accordance with another embodiment of the present invention. Initial aluminum or zinc layer **262** is added to control channel width **260** during rolling. The removable layer **262** is subsequently electroplated with the wall **214** choice of material, for example nickel. After which the aluminum or zinc layer is etched out employing an electroforming technique thus leaving a spiral dewar **250**.

The spiral dewar **250** shown in FIG. **4H** also employs an 55 embossed or folded burner structure **264** that wraps around reformer **32**. FIGS. **41** and **4**J illustrate wash coatings **266** on a wall **268** of burner **30** in accordance with two embodiments of the present invention. For the folded burner structure **264** of FIG. **4I**, a wash coat **266** including the burner catalyst **104** is 60 applied to both sides of wall **268**.

A flat wall **270** suitable for use in spiral dewar **250** is shown in FIG. **4J**. Flat wall **270** includes channels **272** etched or otherwise disposed along its surface. A wash coat **266** including the burner catalyst **104** is then added over the surface of flat wall **270** and channels **272**.

Fuel processors 15 such as that shown in FIG. 4D are very well suited to contain internally generated heat. In one

35

embodiment, burner 30 operates at a temperature greater than about 350 degrees Celsius and the outer side of the housing remains less than about 75 degrees Celsius. This facilitates the use of higher temperature burning fuel sources within burner 30 such as ethanol and propane, for example.

Dewars as shown in FIGS. 4A and 4D may be considered 'multipass' since the incoming air passes over multiple surfaces for convective heat transfer between the warmer surfaces and cooler air. The embodiment in FIG. 4A illustrates a two-pass system where the air passes through two dewar ¹⁰ chambers, while the embodiment in FIG. 4D illustrates an N-pass where N is the number of dewar walls in a given radial direction from the fuel processor center.

FIG. 4E illustrates a cross sectional view of a multipass dewar 300 in accordance with another embodiment of the present invention. Dewar 300 comprises four dewar walls 302a-d that connect to a housing wall 304. Dewar 300 partially contains monolithic structure 100. Dewar wall 302a cooperates with housing wall 304 to enclose monolithic structure 100, which includes burner 30. Dewar wall 302b and housing wall 304 enclose dewar wall 302a and burner 30. Similarly, dewar wall 302c and housing wall 304 enclose dewar wall 302b, while dewar wall 302d and housing wall 304 enclose dewar wall 302c. Dewar walls 302a-d form four volumes for incoming air to pass over warmer walls and ²⁵ receive heat. Air enters dewar inlet port 310 and flows through dewar chamber 308a and into dewar chamber 308b through port 312 after travelling through substantially the whole chamber 308a. Air then serially passes into and through chambers 308c and 308d before entering burner inlet 314.

FIGS. 4F and 4G illustrate a cross section of a fuel processor 15 including a monolithic structure 452 and multipass dewar 450 in accordance with another embodiment of the present invention. Monolithic structure 452 includes multiple reformer chambers 454 that are disposed in a central portion of structure 452. Multiple burner chambers 456 surround and quadrilaterally border the reformer chambers 454. Reformer boiler 458 is arranged within the cross section of burner chambers 456, while burner boiler 460 is arranged in external $_{40}$ portion of the cross section.

Dewar 462 comprises four dewar walls 462a-d. In the cross section shown in FIG. 4F. dewar wall 462a surrounds monolithic structure 452. Dewar wall 462b surrounds and encloses dewar wall 462a. Dewar wall 462c surrounds and encloses 45 dewar wall 462b, while dewar wall 462d surrounds and encloses dewar wall 462c. Dewar walls 462a-d form four dewar volumes for incoming air to pass through and receive heat. As shown in FIG. 4G, air enters dewar inlet port 464 and flows through dewar chamber 468a and into dewar chamber ₅₀ **468***b* after traveling through substantially the whole chamber 468a along the length of monolithic structure 452. Air then serially passes from chamber 468b to chamber 468c and chamber 468d before entering burner inlet 470.

FIG. 5 illustrates a process flow 500 for generating hydro- 55 gen in a fuel processor in accordance with one embodiment of the present invention. The fuel processor comprises a burner, a reformer and a dewar that at least partially contains the burner and reformer. Although the present invention has so far discussed dewars with respect to annular reformer and burner 60 designs described herein, it is also anticipated that dewars described herein are also useful to contain heat in other reformer and burner designs. Many architectures employ a planar reformer disposed on top or below to a planar burner. Micro-channel designs fabricated in silicon commonly 65 employ such stacked planar architectures and would benefit from dewars described herein.

Process flow 500 begins by generating heat in the burner (502). Catalytic burner architectures may include those described above or a micro-channel design on silicon. Further description of a micro-channel fuel processor suitable for use with the present invention is included in commonly owned co-pending patent application entitled "Planar Micro Fuel Processor" naming Ian Kaye as inventor and filed on the same day as this patent application. This application is incorporated by reference for all purposes. A catalyst in the burner facilitates heat generation in the presence of the heating fuel. The burner may also employ an electric burner that includes an resistive heating element that produces heat in response to input current.

Air enters a port for the dewar and passes through a dewar chamber (504). For the dewar of FIG. 4A, burner 30 and dewar 150 share a wall and the air passes through the dewar chamber 156 in a direction that at least partially counters a direction that the air passes through burner chamber 105.

The air is then heated in the dewar chamber using heat generated in the burner (506). Heat travels from burner 30 to dewar chamber 156 via conductive heat transfer. Heat may also travel from a burner to a dewar chamber via convective and/or radiative heat transfer. Once in the dewar chamber, the air is typically heated via convective heat transfer from a wall of the dewar to the air. In one embodiment, the dewar shares a wall with the burner and air in the dewar chamber is heated using heat from the shared wall. Heat from the burner wall may also travel to other walls in the dewar and heat air in the dewar chamber after the heat transfers from the shared wall to another non-shared dewar wall. Heat traveling between the shared wall and non-shared dewar wall transfer by conduction between connected walls or radiation between facing walls.

Process flow 500 then supplies the warmed air to the burner after it has been heated in the dewar chamber (508). Typically, the fuel processor includes an exit to the dewar chamber and an inlet to the burner-along with any intermittent plumbing-that allow the heated air to pass therebetween. For the fuel processor shown in FIG. 2A, space between dewar 150 and burner 30 at the ends of the burner route the air from the dewar to the burner.

The air is then used in the burner for catalytic combustion to generate heat. The generated heat is then transferred from the burner to the reformer (510). In the reformer, the heat is then used in reforming a fuel source to produce hydrogen (512).

The first three elements (502, 504, and 506) of process flow 500 also form a method of managing heat in a fuel processor. In this case, heat generated in the burner (502) passes to air in the dewar (504). The dewar 150 at least partially thermally isolates components internal to the fuel processor housingsuch as the burner-and contains heat within the fuel processor. The dewar thus reduces heat loss from the fuel processor and helps manage the temperature gradient between the burner and outer surfaces of the housing. The dewar may also contain extended and/or multiple dewar chambers through which the air passes and is heated by heat generated in the burner. FIG. 4A illustrates a second dewar chamber 158 formed between the dewar and a housing for the fuel processor. Air passes first through chamber 158 then into dewar chamber 156. FIG. 4D illustrates a spiral dewar including an extended dewar chamber 408. In this case, dewar 250 includes one wall that increasingly provides heat as incoming air nears the burner. FIG. 4E illustrates a dewar 300 including four partially dewar chambers 308 where each dewar chamber heats incoming air in turn as it nears the burner. FIG. 4F illustrates a dewar including four annular and concentric and

40

rectangular dewar chambers 408 that each heat incoming air in turn as it travels to the burner.

While this invention has been described in terms of several preferred embodiments, there are alterations, permutations, and equivalents that fall within the scope of this invention 5 which have been omitted for brevity's sake. For example, although reformer 32 includes chamfered corners as shown in FIG. 3A, the present invention may employ non-chamfered corners in reformer 32. In addition, although the present invention has been described in terms of a monolithic struc- 10 ture 100 that forms the volumetric reformer 32, the present invention is not limited to volumetric reformers disposed in monolithic structures. It is therefore intended that the scope of the invention should be determined with reference to the appended claims.

What is claimed is:

1. A fuel processor for producing hydrogen from a fuel source, the fuel processor comprising:

- a reformer including two or more reformer chambers in a 20 cross section that are configured to receive the fuel source and configured to output hydrogen, each of the two or more reformer chambers including a reformer catalyst that facilitates the production of hydrogen, wherein the two or more reformer chambers are config- $\ ^{25}$ ured such that reactants move through the two or more reformer chambers in series:
- a boiler configured to heat the fuel source before the reformer receives the fuel source;
- a burner including two or more burner chambers in a cross $^{-30}$ section, each of the two or more burner chambers including a burner catalyst, the two or more burner chambers configured to provide heat to the reformer, wherein the two or more burner chambers are configured such that bers in series; and
- a dewar that contains the reformer, contains the burner and includes a set of dewar walls that form a dewar chamber that is configured to receive the fuel source or oxygen before one of the two or more burner chambers receives the fuel source or oxygen,
- wherein the fuel processor includes a monolithic structure having a common material included in walls that define the two or more reformer chambers, the two or more 45 burner chambers and the boiler, wherein the monolithic structure comprises a wall comprising copper that is shared by one of the two or more reformer chambers and one of the two or more burner chambers and the fuel processor is configured to permit conductive thermal 50 communication from the burner catalyst, through the wall, to one of the two or more reformer chambers.

2. The fuel processor of claim 1 wherein the two or more reformer chambers surrounds greater than 50 percent of the two or more burner chamber cross-sectional perimeter.

55 3. The fuel processor of claim 2 wherein the two or more reformer chambers surrounds greater than 75 percent of the two or more burner chamber cross-sectional perimeter.

4. The fuel processor of claim 3 wherein the two or more reformer chambers surrounds greater than 90 percent of the $_{60}$ two or more burner chamber cross-sectional perimeter.

5. The fuel processor of claim 1 wherein the two or more burner chambers bilaterally neighbors the two or more reformer chambers.

6. The fuel processor of claim 5 wherein the two or more 65 burner chambers trilaterally neighbors the two or more reformer chambers.

7. The fuel processor of claim 6 wherein the two or more burner chambers quadrilaterally neighbors the two or more reformer chambers.

8. The fuel processor of claim 1 wherein the two or more burner chambers each has a non-planar cross-sectional shape.

9. The fuel processor of claim 1 wherein the two or more burner chambers each has a non-quadrilateral cross-sectional shape.

10. The fuel processor of claim 1 wherein the dewar includes a radiative layer disposed on an inner wall of the set of dewar walls that improves radiative heat reflectance of the inner wall.

- 11. The fuel processor of claim 1 wherein the dewar is configured such that air passing through the dewar chamber receives heat generated in one of the two or more burner chambers.
- 12. The fuel processor of claim 1 wherein the dewar annularly surrounds the two or more burner chambers in the cross section.

13. The fuel processor of claim 1 wherein the dewar includes a second dewar chamber.

14. The fuel processor of claim 13 wherein the second dewar chamber is configured to receive heat from the two or more burner chambers.

15. The fuel processor of claim 1 wherein the common material comprises a metal.

16. The fuel processor of claim 15 wherein the monolithic structure is formed in a single extrusion or casting.

17. The fuel processor of claim 1 wherein the wall permits conductive thermal communication between the two or more burner chambers and the two or more reformer chambers in orthogonal directions.

18. The fuel processor of claim 17 wherein the boiler is reactants move through the two or more burner cham- ³⁵ configured to receive heat from one of the two or more burner chambers

> 19. The fuel processor of claim 18 further comprising a second boiler that is configured to heat the fuel source before one of the two or more burner chambers receives the fuel source.

> 20. A fuel processor for producing hydrogen from a fuel source, the fuel processor comprising:

- a reformer including two or more reformer chambers in a cross section that are configured to receive the fuel source and configured to output hydrogen, each of the two or more reformer chambers are configured such that reactants move through the two or more reformer chambers in series and including a catalyst that facilitates the production of hydrogen, each of the two or more reformer chambers having a volume greater than about 0.1 cubic centimeters and less than about 50 cubic centimeters, and each of the two or more reformer chambers is characterized by a cross sectional width and a cross sectional height that is greater than one-third the cross sectional width;
- a boiler configured to heat the fuel source before the reformer receives the fuel source;
- a burner including two or more burner chambers in a cross section, each of the burner chambers including a burner catalyst, each of the two or more burner chambers are configured to provide heat to the reformer, wherein the two or more burner chambers are configured such that reactants move through the two or more burner chambers in series:
- a dewar that contains the reformer, contains the burner and includes a set of dewar walls that form a dewar chamber

35

that is configured to receive the fuel source or oxygen before one of the two or more burner chambers receives the fuel source or oxygen,

wherein the fuel processor includes a monolithic structure comprising copper that forms a wall of one of the two or 5 more reformer chambers, a wall of one of the two or more burner chambers, and a wall of the boiler.

21. The fuel processor of claim **20** wherein the two or more reformer chambers each are substantially non-quadrilateral in cross section.

22. The fuel processor of claim **20** wherein the two or more reformer chambers each includes a substantially elliptical cross section.

23. The fuel processor of claim 22 wherein the two or more reformer chambers each include a substantially circular cross 15 section.

24. The fuel processor of claim 20 wherein the two or more reformer chambers each comprises a volume between about 0.5 cubic centimeters and about 2.5 cubic centimeters.

25. The fuel processor of claim **20** wherein the cross sec- 20 tional height is greater than one-half the cross sectional width.

26. The fuel processor of claim 20 wherein the cross sectional height is greater than the cross sectional width.

27. The fuel processor of claim **20** wherein the two or more burner chambers each has a non-quadrilateral cross-sectional 25 shape.

28. The fuel processor of claim **27** wherein the two or more burner chambers bilaterally neighbors the two or more reformer chambers.

29. The fuel processor of claim **28** wherein the two or more 30 burner chambers collectively quadrilaterally neighbor the two or more reformer chambers.

30. The fuel processor of claim **20** wherein the two or more reformer chambers surrounds greater than 90 percent of the two or more burner chamber perimeter.

31. The fuel processor of claim **20** wherein the boiler is configured to receive heat from at least one of the two or more burner chambers.

32. The fuel processor of claim **20** wherein the fuel processor includes a monolithic structure having a common 40 material included in walls that define the two or more reformer chambers, the two or more burner chambers and the boiler.

33. The fuel processor of claim **32** wherein the common material comprises a metal.

34. The fuel processor of claim **33** wherein the monolithic structure is formed in a single extrusion.

35. The fuel processor of claim **20** wherein the catalyst comprises pellets.

36. The fuel processor of claim **35** wherein the pellets have a diameter ranging from about 300 microns to about 1500 microns.

37. A fuel processor for producing hydrogen from a fuel source, the fuel processor comprising:

- a reformer including two or more reformer chambers in a cross section that are configured to receive the fuel source and configured to output hydrogen, each of the two or more reformer chambers including a reformer catalyst that facilitates the production of hydrogen, wherein the two or more reformer chambers are configured such that reactants move through the two or more reformer chambers in series;
- a boiler configured to heat the fuel source before the reformer receives the fuel source;
- a burner, including two or more burner chambers in a cross section, each of the two or more burner chambers include a burner catalyst, each of the two or more burner chambers configured to provide heat to the reformer;
- a dewar that contains the reformer, contains the burner and includes a set of dewar walls that form a dewar chamber that is configured to receive the fuel source or oxygen before one of the two or more burner chambers receives the fuel source or oxygen,
- wherein the fuel processor includes a monolithic structure comprising copper that forms a wall of at least one of the two or more reformer chambers, a wall of at least one of the two or more burner chambers, and a wall of the boiler.

38. The fuel processor of claim **20** wherein the two or more reformer chambers surrounds greater than 50 percent of the two or more burner chamber cross-sectional perimeter.

39. The fuel processor of claim **20** wherein the two or more reformer chambers surrounds greater than 75 percent of the two or more burner chamber cross-sectional perimeter.

40. The fuel processor of claim **37** wherein the two or more burner chambers are configured such that reactants move through the two or more burner chambers in series.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE CERTIFICATE OF CORRECTION

PATENT NO. : 7,604,673 B2 APPLICATION NO. : 10/877044 DATED : October 20, 2009 INVENTOR(S) : Kaye et al. Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

On the Title Page:

The first or sole Notice should read --

Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 929 days.

Signed and Sealed this

Fifth Day of October, 2010

Jand J. g/pos

David J. Kappos Director of the United States Patent and Trademark Office