Decision Diagnostics Corp.

ANNUAL REPORT REPORT FOR OTC PINK

Supplemental Disclosures

Quarterly Report for Year Ended

December 31, 2017

Trading Symbol: **DECN**
CUSIP Number: **243443 108**

Decision Diagnostics Corp.

OTC Pink Basic Disclosure Guidelines

1) Name of the issuer and its predecessors (if any)

In answering this item, please also provide any names used by predecessor entities in the past five years and the dates of the name changes.

DECISION DIAGNOSTICS CORP. (11/25/2011-present)
INSTACARE CORP. (through 11/25/2011)

2) Address of the issuer's principal executive offices

Company Headquarters
Address 1: 2660 TOWNSGATE ROAD
Address 2: SUITE 300
Address 3: WESTLAKE VILLAGE, CA 91361
Phone: 805-446-1973
Email: info@decisiondiagnostics.com
Website(s): www.decisiondiagnostics.com

IR Contact N/A

3) Security Information

Trading Symbol: DECN
Exact title and class of securities outstanding: COMMON
CUSIP: **243443 108**
Par or Stated Value: $0.001
Total shares authorized: 494,995,000 as of: 12/31/2017
Total shares outstanding: 110,241,640 as of: 12/31/2017

Additional class of securities (if necessary):
Trading Symbol: N/A
Exact title and class of securities outstanding: PREFERRED
CUSIP: N/A
Par or Stated Value: $0.001
Total shares authorized: 3,738,500 as of: 12/31/2017
Total shares outstanding: N/A as of: 12/31/2017

Additional class of securities (if necessary):
Trading Symbol: N/A
Exact title and class of securities outstanding: PREFERRED SERIES "B"
CUSIP: N/A
Par or Stated Value: $0.001
Total shares authorized: 2,500 as of: 12/31/2017
Total shares outstanding: 1,000 as of: 12/31/2017*

(*) The company rescinded all outstanding Preferred B shares during 1Q 2018 resulting from criminal issues surrounding the sole Preferred B shareholder.

Additional class of securities (if necessary):
Trading Symbol: N/A
Exact title and class of securities outstanding: PREFERRED SERIES "C"
CUSIP: N/A
Par or Stated Value: $0.001
Total shares authorized: 10,000 as of: 12/31/2017
Total shares outstanding: 6,235 as of: 12/31/2017

Additional class of securities (if necessary):
Trading Symbol: N/A
Exact title and class of securities outstanding: PREFERRED SERIES "D"
CUSIP: N/A
Par or Stated Value: $0.001
Total shares authorized: 1,250 as of: 12/31/2017
Total shares outstanding: 40 as of: 12/31/2017

Additional class of securities (if necessary):
Trading Symbol: N/A
Exact title and class of securities outstanding: PREFERRED SERIES "E"
CUSIP: N/A
Par or Stated Value: $0.001
Total shares authorized: 1,250,000 as of: 12/31/2017
Total shares outstanding: 813,240 as of: 12/31/2017

Transfer Agent
Name: ACTION STOCK TRANSFER CORP.
Address 1: 2469 E. FORT UNION BLVD.
Address 2: SUITE 214
Address 3: SALT LAKE CITY, UT 84121
Phone: 801-274-1088

Is the Transfer Agent registered under the Exchange Act?* Yes: XX No: ☐

*To be included in the OTC Pink Current Information tier, the transfer agent must be registered under the Exchange Act.

In April 2015 the company completed voluntary disclosure, periodic financial, and management's discussion and analysis filings (postings) with OTCMarkets, for the purposes of becoming a current voluntary filer. The company's filings were reviewed and the company was granted current filer status with OTCMarkets on April 21, 2015. The company had to repeat this process in August 2016.

List any restrictions on the transfer of security:

None

Describe any trading suspension orders issued by the SEC in the past 12 months.

None

List any stock split, stock dividend, recapitalization, merger, acquisition, spin-off, or reorganization either currently anticipated or that occurred within the past 12 months:

1:14 reverse stock split of $0.001 par value common stock effective 11/25/2011

4) Issuance History

List below any events, in chronological order, that resulted in changes in total shares outstanding by the issuer in the past two fiscal years and any interim period. The list shall include all offerings of equity securities, including debt convertible into equity securities, whether private or public, and all shares or any other securities or options to acquire such securities issued for services, describing (1) the securities, (2) the persons or entities to whom such securities were issued and (3) the services provided by such persons or entities. The list shall indicate:

COMMON STOCK

Date	Description	Change in Shares	Running Total
12/1/2011	1 for 14 Reverse Split	8,461,032	8,461,032
12/19/2011	New Issuance-Kimberly Binder	75	8,461,107
12/19/2011	New Issuance-Patrick DiParini	200	8,461,307
12/30/2011	10% Stock Dividend	846,669	9,307,976
1/3/2012	DTC Rounding shares	(42)	9,307,934
1/4/2012	New Issuance-Positive Revolution Inc-S-8	100,000	9,407,934
1/11/2012	Converted to Common-Alpha Credit	294,000	9,701,934
1/18/2012	New Issuance-Debt Conv. Andrew Edenbaum	53,354	9,755,288
1/23/2012	DTC Rounding shares	25	9,755,313
3/5/2012	New Issuance-JFS Investments Inc	60,000	9,815,313
3/5/2012	New Issuance-Garden State Securities	60,000	9,875,313
3/5/2012	New Issuance-Excell Advisors	30,000	9,905,313
3/5/2012	Return to Treasury-Positive Revolution	(100,000)	9,805,313
3/5/2012	New Issuance-TPC Holdings Group-ESOP-06	300,000	10,105,313
3/5/2012	New Issuance-Cadence Consulting-ESOP-06	50,000	10,155,313
3/30/2012	New Issuance-Alpha Credit Resources	238	10,155,551
6/27/2012	New Issuance-Rocio C Carazas-ESOP-06	375,000	10,530,551
6/27/2012	New Issuance-Marjolein Imfeld-ESOP-06	375,000	10,905,551
9/26/2012	Converted to Common-Centurion	172,200	11,077,751
10/9/2012	New Issuance-Aubyn Inc-ESOP-06	400,000	11,477,751
11/8/2012	Return to Treasury-Aubyn Inc-ESOP-06	(200,000)	11,277,751
11/8/2012	New Issuance-Mayer & Assoc. Esop-04	650,000	11,927,751
11/8/2012	New Issuance-Mayer & Associates	200,000	12,127,751
11/8/2012	New Issuance-Curing Capital Inc	400,000	12,527,751
11/13/2012	Converted to Common-Centurion	182,000	12,709,751
11/13/2012	New Issuance-Econ Corporate Services	50,000	12,759,751
11/13/2012	New Issuance-Call Van Zant-ESOP-06	100,000	12,859,751
11/13/2012	New Issuance-Darren Bankstead-ESOP-06	50,000	12,909,751
11/13/2012	New Issuance-Axiom Financial Inc	200,000	13,109,751
12/21/2012	Cancellation-Mayer & Associates LLC	(200,000)	12,909,751
12/21/2012	New Issuance-Mayer & Associates LLC	1,000,000	13,909,751
1/7/2013	New Issuance-Mayer & Associates LLC	50,000	13,959,751
1/7/2013	Converted to Common-Apex Clearing	210,000	14,169,751
1/7/2013	Converted to Common-Apex Clearing	236,600	14,406,351
2/15/2013	New Issuance-TPC Holdings Group-ESOP	1,325,000	15,731,351
2/15/2013	New Issuance-Envisionte LLC-ESOP	700,000	16,431,351
2/15/2013	New Issuance-Bridgeview Capital Group ESOP	700,000	17,131,351
2/15/2013	New Issuance-Cadence Holdings LLC ESOP	275,000	17,406,351
2/15/2013	New Issuance-AAC Group LLC ESOP	600,000	18,006,351
2/15/2013	New Issuance-Cadence Holdings LLC ESOP	150,000	18,156,351
2/15/2013	New Issuance-St Andrews Inc	1,000,000	19,156,351
2/15/2013	New Issuance-Alan Binder ESOP	100,000	19,256,351
2/15/2013	New Issuance-Dale Richter ESOP	100,000	19,356,351

2/15/2013	New Issuance-Kimberly Binder ESOP	50,000	19,406,351
2/15/2013	New Issuance-Maria Luz Johnson-ESOP	25,000	19,431,351
2/18/2013	Converted to Common-Apex Clearing	324,800	19,756,151
2/22/2013	New Issuance-Robert Herskowitz ESOP	500,000	20,256,151
2/22/2013	New Issuance-Jeff Whitelaw	125,000	20,381,151
2/22/2013	New Issuance-Brent England	75,000	20,456,151
5/9/2013	Converted to Common-Apex Clearing	868,000	21,324,151
5/10/2013	Cancellation-Robert Herskowitz ESOP	(500,000)	20,824,151
5/10/2013	Cancellation-St. Andrews	(1,000,000)	19,824,151
5/10/2013	New Issuance-Chase Financing Inc ESOP	350,000	20,174,151
5/10/2013	New Issuance-Mayer & Associates LLC ESOP	1,000,000	21,174,151
8/7/2013	New Issuance-St Andrews Inc ESOP	500,000	21,674,151
8/15/2013	New Issuance-Robert Herskowitz ESOP	25,000	21,699,151
8/27/2013	Cancellation-Curring Capital	(200,000)	21,499,151
8/27/2013	Cancellation-ACC Group ESOP	(600,000)	20,899,151
8/27/2013	New Issuance-Benjamin Mayer ESOP	950,000	21,849,151
9/20/2013	New Issuance-SLCC Partners LLC	1,000,000	22,849,151
9/20/2013	New Issuance-Envisionte LLC-ESOP	500,000	23,349,151
9/20/2013	New Issuance-Thomas Hanson-ESOP	250,000	23,599,151
9/20/2013	New Issuance-Envisionte LLC-ESOP	250,000	23,849,151
10/2/2013	New Issuance-Joanne Broeders-ESOP	235,300	24,084,451
10/2/2013	Cancellation-Alan Binder ESOP	(100,000)	23,984,451
10/2/2013	New Issuance-Kimberly Binder	100,000	24,084,451
10/2/2013	Converted to Common-COR Clearing	1,078,000	25,162,451
10/28/2013	Converted to Common-Michael Belcher	350,000	25,512,451
10/28/2013	New Issuance	2,798,728	28,311,179
10/30/2013	New Issuance-Benjamin Mayer ESOP	100,000	28,411,179
10/30/2013	New Issuance-Benjamin Mayer	300,000	28,711,179
10/30/2013	New Issuance	166,365	28,877,544
11/11/2013	Conversion-Centurion Credit	980,000	29,857,544
11/11/2013	New Issuance-Benjamin Mayer ESOP	500,000	30,357,544
11/11/2013	New Issuance	125,000	30,482,544
12/4/2013	Conversion-Centurion Credit	1,220,800	31,703,344
12/23/2013	New Issuance-Mark Herskowitz ESOP	175,000	31,878,344
12/23/2013	New Issuance-Benjamin Mayer ESOP	600,000	32,478,344
12/23/2013	New Issuance	1,200,548	33,678,892
1/2/2014	New Issuance	2,709,678	36,388,570
1/15/2014	New Issuance	748,720	37,137,290
1/15/2014	New Issuance	267,105	37,404,395
2/18/2014	Conversion-Alpha Credit	611,940	38,016,335
2/18/2014	Conversion-Michael Belcher	350,000	38,366,335
2/19/2014	Conversion-Mayer & Associates	798,000	39,164,335
3/28/2014	Conversion-Alpha Credit	523,740	39,688,075
3/28/2014	New Issuance	400,000	40,088,075
6/3/2014	Conversion-Alpha Credit	499,996	40,588,071
6/4/2014	Conversion-Mayer & Associates	1,115,660	41,703,731
8/14/2014	Conversion-Alpha Credit	245,000	41,948,731
8/15/2014	Conversion-Mayer & Associates	550,000	42,498,731
9/9/2014	Conversion-Mayer & Associates	775,000	43,273,731
10/28/2014	Conversion	675,010	43,948,741
1/21/2015	New Issuance	1,875,000	45,823,741
1/28/2015	New Issuance	850,000	46,673,741

2/23/2015	Conversion-Alpha Credit	705,124	47,378,865
5/11/2015	New Issuance-Momona Capital	235,000	47,613,865
5/12/2015	Conversion-Mayer & Associates	950,040	48,563,905
5/12/2015	New Issuance-Robert Herskowiz	950,000	49,513,905
5/21/2015	New Issuance-Momona Capital	235,000	49,748,905
6/1/2015	New Issuance-Chase Financing 401K	533,334	50,282,239
6/8/2015	New Issuance-Momona Capital	437,250	50,719,489
6/8/2015	New Issuance-St Andrews	350,000	51,069,489
6/29/2015	New Issuance-Alpha Capital Anstalt	384,537	51,454,026
7/27/2015	New Issuance-Alpha Capital Anstalt	387,907	51,841,933
8/24/2015	New Issuance-Alpha Capital Anstalt	313,022	52,154,955
9/16/2015	Conversion-Mayer & Associates	1,890,000	54,044,955
9/16/2015	Conversion-Robert Herskowitz	1,400,000	55,444,955
10/27/2015	New Issuance-Alpha Capital Anstalt	479,489	55,924,444
12/2/2015	New Issuance-Alpha Capital Anstalt	950,545	56,874,989
12/15/2015	New Issuance-Alpha Capital Anstalt	950,545	57,825,534
12/21/2015	New Issuance-Alpha Capital Anstalt	956,950	58,782,484
2/2/2016	New Issuance-Alpha Capital Anstalt	970,980	59,753,464
2/17/2016	New Issuance-Alpha Capital Anstalt	1,614,248	61,367,712
2/25/2016	New Issuance-Robert Herskowiz	750,000	62,117,712
3/21/2016	New Issuance-Paradigm Capital Holdings	1,400,000	63,517,712
3/21/2016	New Issuance-Robert Herskowitz	200,000	63,717,712
3/29/2016	New Issuance-Alpha Capital Anstalt	404,630	64,122,342
3/29/2016	New Issuance-James J Loures	500,000	64,622,342
4/13/2016	New Issuance-Robert Herskowitz	280,000	64,902,342
4/13/2016	New Issuance-Robert Herskowitz	280,000	65,182,342
4/13/2016	New Issuance-Robert Herskowitz 2011 Irv TR	140,000	65,322,342
4/13/2016	New Issuance-Chase Financial	148,160	65,470,502
4/13/2016	New Issuance-Mark Herskowittz	185,195	65,655,697
4/13/2016	New Issuance-Andrew Schoenzeit	37,040	65,692,737
4/13/2016	New Issuance-Robert Herskowitz 2011 Irv TR	431,376	66,124,113
4/26/2016	New Issuance-LICGO Partners	1,837,500	67,961,613
4/26/2016	Conversion-Mayer & Associates	200,200	68,161,813
5/2/2016	New Issuance-Robert Herskowitz	472,106	68,633,919
5/5/2016	New Issuance-Alpha Capital Anstalt	998,099	69,632,018
5/17/2016	New Issuance-Alpha Capital Anstalt	422,669	70,054,687
5/17/2016	New Issuance-Navesink	625,000	70,679,687
5/18/2016	New Issuance-LICGO Partners	525,000	71,204,687
5/18/2016	Conversion-Mayer & Associates	220,000	71,424,687
6/1/2016	New Issuance-Alpha Capital Anstalt	814,314	72,239,001
6/6/2016	New Issuance-Mark Herskowitz	1,000,000	73,239,001
6/6/2016	New Issuance-Chase Financing Inc Profit Sh.	1,050,000	74,289,001
6/6/2016	New Issuance-Robert Herskowitz	280,000	74,569,001
6/6/2016	New Issuance-Robert Herskowitz 2011 Irv TR	70,000	74,639,001
6/8/2016	New Issuance-Alpha Capital Anstalt	484,148	75,123,149
6/27/2016	New Issuance-Navesink	625,000	75,748,149
7/18/2016	New Issuance-Cadence Holdings LLC	100,000	75,848,149
7/18/2016	New Issuance-TPC Holdings Group	150,000	75,998,149
7/21/2016	New Issuance-Robert Herskowitz	700,000	76,698,149
7/21/2016	New Issuance-Robert Herskowitz 2011 Irv TR	70,000	76,768,149
7/21/2016	New Issuance-Chase Financial	945,000	77,713,149
8/2/2016	New Issuance-Navesink	625,000	78,338,149

8/29/2016	New Issuance-Alpha Capital Anstalt	954,925	79,293,074
9/7/2016	New Issuance-Chase Financial	945,000	80,238,074
9/19/2016	New Issuance-Alpha Capital Anstalt	521,784	80,759,858
9/19/2016	New Issuance-Mark Herskowitz	805,147	81,565,005
9/19/2016	New Issuance-Marc Berger	400,000	81,965,005
11/21/2016	New Issuance-Alpha Capital Anstalt	957,485	82,922,490
12/6/2016	New Issuance-Alpha Capital Anstalt	962,118	83,884,608
12/12/2016	New Issuance-LICGO Partners	755,300	84,639,908
1/9/2017	New Issuance-Alpha Capital Anstalt	971,074	85,610,982
1/9/2017	New Issuance-Mark Herskowitz	400,000	86,010,982
3/1/2017	New Issuance-Alpha Capital Anstalt	989,425	87,000,407
3/3/2017	New Issuance-Chase Financial	1,400,000	88,400,407
3/3/2017	New Issuance-Robert Herskowitz	560,000	88,960,407
3/3/2017	New Issuance-R Herskowitz 2011 Irrv. TR	140,000	89,100,407
3/10/2017	Issuance-Mark Herskowitz	400,000	89,500,407
3/21/2017	New Issuance-Alpha Capital Anstalt	355,803	89,856,210
4/19/2017	New Issuance-Paradigm Capital Holdings	400,000	90,256,210
5/10/2017	New Issuance-Navesink	625,000	90,881,210
5/17/2017	New Issuance-OmniVance Advisors LLC	100,000	90,981,210
6/19/2017	New Issuance-Alpha Capital Anstalt	1,096,312	92,077,522
7/11/2017	New Issuance-Robert Herskowitz	1,400,000	93,477,522
7/11/2017	New Issuance-Chase Financial	1,400,000	94,877,522
7/24/2017	New Issuance-Navesink	625,000	95,502,522
7/24/2017	New Issuance-Paradigm Capital Holdings	1,475,000	96,977,522
8/1/2017	New Issuance-Mark Herskowitz	350,000	97,327,522
8/7/2017	New Issuance-Alpha Capital Anstalt	981,067	98,308,589
8/21/2017	New Issuance-Alpha Capital Anstalt	971,043	99,279,632
8/24/2017	New Issuance-R Herskowitz 2011 Irrv. TR	700,000	99,979,632
9/5/2017	New Issuance-Mark Herskowitz	350,000	100,329,632
9/20/2017	New Issuance-Alpha Capital Anstalt	952,043	101,281,675
10/3/2017	New Issuance-Alpha Capital Anstalt	987,640	102,269,315
10/23/2017	New Issuance-Alpha Capital Anstalt	991,943	103,261,258
11/6/2017	New Issuance-Mark Herskowitz	500,000	103,761,258
11/6/2017	New Issuance-Alpha Capital Anstalt	2,878,058	106,639,316
12/4/2017	New Issuance-Alpha Capital Anstalt	1,502,294	108,141,610
12/6/2017	New Issuance-Chase Financing Inc	700,000	108,841,610
12/12/2017	New Issuance-Scott J Weiner	1,000,000	109,841,610
12/19/2017	New Issuance-Robert Herskowitz	1,400,000	111,241,610
12/31/2017	Cancellation-Scott J Weiner	(1,000,000)	110,241,610

PREFERRED B STOCK*

Date	Description	Change in Shares	Running Total
3/23/2011	New Issuance-Centurion Credit Resources	1,000	1,000

PREFERRED C STOCK

Date	Description	Change in Shares	Running Total
1/4/2012	New Issuance-Michael Belcher	1,250	1,250
8/27/2013	New Issuance-Lathrop Gage LLC	1,500	2,750
10/28/2013	Conversion-Michael Belcher	(70)	2,680
2/18/2014	Conversion-Michael Belcher	(70)	2,610

12/30/2015	New Issuance-Navesink Device Initiatives	1,475	4,085
3/21/2016	New Issuance-Paradigm Capital	800	4,885
4/26/2016	New Issuance-LICGO Partners	1,050	5,935
4/26/2016	New Issuance-Paradigm Capital	325	6,260
5/17/2016	Conversion-Navesink Device	(125)	6,135
5/18/2016	New Issuance-LICGO Partners	300	6,435
5/18/2016	New Issuance-Paradigm Capital	50	6,485
6/27/2016	Conversion-Navesink Device	(125)	6,360
8/2/2016	Conversion-Navesink Device	(125)	6,235
4/19/2017	Conversion-Paradigm Capital	(80)	6,155
4/19/2017	New Issuance-LICGO Partners	157	6,312
5/10/2017	Conversion-Navesink Device	(125)	6,187
7/24/2017	Conversion-Navesink Device	(125)	6,062
7/24/2017	Conversion-Paradigm Capital	(295)	5,767
7/25/2017	New Issuance-LICGO Partners	196	5,963
9/28/2017	New Issuance-Gerald Hickson	300	6,263
10/23/2017	New Issuance-LICGO Partners	210	6,473

PREFERRED D STOCK

Date	Description	Change in Shares	Running Total
12/31/2017	New Issuance-Sovereign Partners	40	40

PREFERRED E STOCK

Date	Description	Change in Shares	Running Total
8/1/2008	New Issuance-Centurion Credit	14,900	14,900
11/5/2008	New Issuance-Centurion Credit	21,225	36,125
12/16/2008	New Issuance-Centurion Credit	30,785	66,910
1/15/2009	New Issuance-Centurion Credit	40,000	106,910
3/31/2009	New Issuance-Centurion Credit	23,000	129,910
3/31/2009	New Issuance-Centurion Credit	19,000	148,910
4/1/2009	Converted to Common	(14,900)	134,010
5/13/2009	New Issuance-Centurion Credit	17,800	151,810
6/2/2009	New Issuance-Centurion Credit	25,000	176,810
7/8/2009	New Issuance-Centurion Credit	25,000	201,810
8/13/2009	New Issuance-Centurion Credit	13,000	214,810
9/11/2009	New Issuance-Centurion Credit	12,600	227,410
10/7/2009	New Issuance-Centurion Credit	20,000	247,410
11/4/2009	New Issuance-Centurion Credit	16,700	264,110
11/18/2009	New Issuance-Centurion Credit	60,000	324,110
11/20/2009	Converted to Common	(92,010)	232,100
11/23/2009	Converted to Common	(59,800)	172,300
12/7/2009	Converted to Common	(25,000)	147,300
12/8/2009	New Issuance-Centurion Credit	720,000	867,300
1/20/2010	Converted to Common	(25,000)	842,300
2/16/2010	Converted to Common	(13,000)	829,300
3/17/2010	Converted to Common	(12,600)	816,700
4/16/2010	Converted to Common	(20,000)	796,700
5/25/2010	Converted to Common	(16,700)	780,000
6/4/2010	Converted to Common	(60,000)	720,000
7/19/2010	Converted to Common	(10,000)	710,000
8/4/2010	New Issuance-Centurion Credit	200,000	910,000
1/26/2011	Converted to Common	(54,500)	855,500

3/8/2011	New Issuance-Centurion Credit	240,000	1,095,500
5/17/2011	Converted to Common	(135,200)	960,300
5/17/2011	New Issuance-Centurion Credit	135,000	1,095,300
1/11/2012	Converted to Common	(21,000)	1,074,300
3/30/2012	New Issuance-Alpha Credit Resources	124,700	1,199,000
9/26/2012	Converted to Common	(12,300)	1,186,700
11/13/2012	Converted to Common	(13,000)	1,173,700
1/7/2013	Converted to Common	(15,000)	1,158,700
1/7/2013	Converted to Common	(16,900)	1,141,800
2/18/2013	Converted to Common	(23,200)	1,118,600
5/9/2013	Converted to Common	(62,000)	1,056,600
10/2/2013	Converted to Common	(77,000)	979,600
11/11/2013	Conversion-Centurion Credit	(70,000)	909,600
12/4/2013	Conversion-Centurion Credit	(87,200)	822,400
1/15/2014	Conversion-Alpha Credit	(53,480)	768,920
2/18/2014	New Issuance-Mayer & Associates	125,000	893,920
2/18/2014	Conversion-Alpha Credit	(43,710)	850,210
2/19/2014	Conversion-Mayer & Associates	(57,000)	793,210
3/28/2014	Conversion-Alpha Credit	(37,400)	755,810
6/3/2014	Conversion-Alpha Credit	(35,714)	720,096
6/4/2014	Conversion-Mayer & Associates	(79,690)	640,406
8/14/2014	Conversion-Alpha Credit	(17,500)	622,906
8/15/2014	Conversion-Mayer & Associates	(39,285)	583,621
9/9/2014	Conversion-Mayer & Associates	(55,357)	528,264
10/28/2014	Conversion-Mayer & Associates	(30,358)	497,906
1/21/2015	New Issuance-Robert Herskowitz	100,000	597,906
1/21/2015	New Issuance-Mayer & Associates	135,000	732,906
1/21/2015	New Issuance-Alpha Credit Resources	67,860	800,766
2/23/2015	New Issuance-Alpha Credit Resources	(50,366)	750,400
5/12/2015	Conversion-Mayer & Associates	(67,860)	682,540
5/12/2015	New Issuance-Robert Herskowitz	30,000	712,540
7/27/2015	New Issuance-Chase Financing	75,000	787,540
9/16/2015	Conversion-Mayer & Associates	(135,000)	652,540
9/16/2015	Conversion-Robert Herskowitz	(100,000)	552,540
9/16/2015	New Issuance-Chase Financing	135,000	687,540
2/25/2016	New Issuance-Robert Herskowitz	100,000	787,540
3/21/2016	New Issuance-Mayer & Associates	14,300	801,840
4/26/2016	Conversion-Mayer & Associates	(14,300)	787,540
6/6/2016	New Issuance-Mark Herskowitz 401K Trust	100,000	873,240
6/6/2016	New Issuance-Chase Financing Inc Profit Sh.	35,000	908,240
6/6/2016	New Issuance-Chase Financing	100,000	1,008,240
6/6/2016	Conversion-Chase Financing Inc Profit Sh.	(75,000)	933,240
7/21/2016	Conversion-Chase Financing Inc	(67,500)	865,740
7/21/2016	Conversion-Robert Herskowitz	(30,000)	835,740
9/7/2016	Conversion-Chase Financing Inc	(67,500)	768,240
9/19/2016	New Issuance-Chase Financing Inc Profit Sh.	75,000	843,240
1/9/2017	New Issuance-Chase Financing Inc Profit Sh.	105,000	948,240
3/3/2017	Cancellation	(105,000)	843,240
3/3/2017	New Issuance-Chase Financing	50,000	893,240
3/3/2017	New Issuance-Chase Financing Inc Profit Sh.	70,000	963,240
3/3/2017	Conversion-Chase Financing	(100,000)	863,240
5/17/2017	New Issuance-Chase Financing	100,000	963,240

7/11/2017	Conversion-Robert Herskowitz	(100,000)	863,240
7/11/2017	Conversion-Chase Financing	(100,000)	763,240
8/24/2017	New Issuance-Chase Financing	50,000	813,240
8/24/2017	New Issuance-Chase Financing Inc Profit Sh.	50,000	863,240
8/24/2017	New Issuance-Chase Financing	50,000	827,540
8/24/2017	New Issuance-Chase Financing Inc Profit Sh.	50,000	877,540
12/6/2017	New Issuance-Chase Financing	(50,000)	827,540
12/12/2017	New Issuance-Robert Herskowitz	100,000	927,540
12/19/2017	Conversion-Robert Herskowitz	(100,000)	827,540
12/31/2017	Cancellation-Benjamin Mayer (2016 Issuance)	(14,300)	813,240

A. Whether the certificates or other documents that evidence the shares contain a legend (1) stating that the shares have not been registered under the Securities Act and (2) setting forth or referring to the restrictions on transferability and sale of the shares under the Securities Act.

See above

5) Financial Statements

SEE FINANCIAL STATEMENTS ATTACHED TO THIS DISCLOSURE STATEMENT

6) Describe the Issuer's Business, Products and Services

Overview

Decision Diagnostics Corp. is a worldwide prescription and non-prescription diagnostics and home testing products distributor and the manufacturer of the Genstrip 50 (discontinued in November 2016) and GenUltimate! glucose test strips, both Class II medical devices for at-home use for the measurement of glucose. The company also has its GenSure! glucose test strip, a product for off-shore sales which is complete and available for sales, but will primarily be sold as an International private label market entry. In addition, the company's GenChoice! glucose test strip is in patient clinical trials, and its GenPrecis! test strip and Precise meter have just begun patient testing (first level clinical trials). As an off-shore product GenSure! is currently in registration and is seeking a CE mark in the EU. It was launched in 4Q 2017. We have identified International distributors for this product. At the conclusion of the respective clinical trials, the GenChoice! And GenPrecis! products will be registered in the EU and applications for 510K pre-market will be filed with the FDA. The company has contracted with the expert organization that will write the 510K document and prosecute these documents along with the company.

The U.S. FDA, in a manner similar to prescription drugs, regulates diagnostic test kits and at-home patient testing products a similar but somewhat streamlined process, to the regulation of prescription medicine. The regulatory standard used for the Genstrip 50 was the 510k pre-market and post-market processes. The same process will be used for the GenChoice! And GenPrecis! products beginning with the 510k approval with the FDA during the latter part of 2018. Both the GenChoice! And GenPrecis! products will be sold internationally while the U.S. FDA 510k applications are pending.

Previous to this change in business model, from 2005 and until 2013, the company contracted with independent pharmacies for use of their prescription drug distribution licenses. At that time the company made market and sold brand name over the counter pharmaceutical items with a concentration in legacy diabetic test strips. The brand name products we distributed, for the most part, did not require a doctor's prescription for anything other than insurance benefit compliance. Our previous business model worked well in the previous regulated environment, although the financial benefits were stressed by major changes made to the Federal Medicare plan that have led to substantially lower rates of reimbursement and ultimately an unprofitable business model.

Our Current Business Foundation

Our subsidiaries, Pharma Tech Solutions, Inc., PDA Services, Inc. and PharmaTech Sensor Development Corp. operate in several healthcare products channels. In addition our subsidiary Decision IT Corp. engages in the acquisition and holding of Intellectual Property including Patents and Trademarks and specialty manufacturing equipment acquired for our Korean contract

manufacturer of our GenUltimate! and our in development GenSure! and GenChoice! products. Our newest subsidiary Pharmatech Sensor Development Corp. manages our investment in specialty manufacturing machinery and testing laboratories, as well as an inventory credit line to finance inventory purchases of our Genstrip 50 and GenUltimate! products. This credit line will be expanded for the management of our GenChoice! And GenPrecis! products in 2018. The company has discontinued its GenStrip 50 product and ended the selling of the last of the inventory in November 2016.

In March 2017 the company was approached by its Korean partner, The Bio Co., Ltd to design and fund a new product which the company calls GenPrecis!. This product represents a major improvement in diabetic glucose monitoring. The GenPrecis! system will be the first of its kind +/- 9% system. Current ISO (2015) and FDA (2014) guidelines call for glucose monitoring systems to meet a +/- 15% standard, whereby the meter and strip must be within +/- 15% of a reference method in repeated samplings 95% of the time. GenUltimate! and GenChoice! are +/- 15% test strips, but in each case 97+% of the time in repeated samplings. GenPrecis! is designed to meet the written standards of the ISO and FDA at +/- 9%, 97% of the time – effectively setting a new standard. The company has been funding the development of this system product in 2017, as well as a test strip only derivative version for use with a legacy meter sold overseas. The system product will be ready for testing in June 2018 and will be registered for International sale at that time. However, the natural market for this product will be the U.S, and Canada where precision standards are higher for new products.

As of this writing, neither GenSure! nor GenPrecis! is available for sale or distribution in the U.S. or Puerto Rico.

From time to time, when economic conditions warrant and given market conditions, we distribute other brand name prescription and non-prescription diagnostics products, as well as several lines of ostomy, wound care and post-surgery medical products, although these healthcare channels have also undergone two major market changes and disruptions since July 2013 and we have determined that we will maintain our contacts but in 2017 we refrained from competing. Our main product was the Genstrip 50 and its successor brand the GenUltimate!, both of improved performance and design improvements and a rebranding and development (from scratch) of the original Shasta Technologies Genstrip. Both of these glucose test strips are of our manufacture. We maintain FDA registered contract manufacturers in Pennsylvania and South Korea,. We ended our association with the contract manufacturer in Pennsylvania as of March 31, 2017. The original GenStrip was cleared for market by the FDA on November 30, 2012. By virtue of our written agreements with Shasta in 2011, we were granted an irrevocable license to prosecute their 510k application with the U.S. FDA, and we succeeded. This was no small feat. We introduced the original Genstrip in March 2013. We then acquired Genstrip from Shasta Technologies LLC on March 20, 2014 and in late June 2014 we made the first branding changes. We began work on the GenUltimate! product in July 2015 and introduced this improved test strip (vs. our GenStrip) in April 2016. The original Shasta Genstrip and our Genstrip 50 have been discontinued.

Historical Construct

Shasta Technologies LLC, the original specifications provider of GenStrip, had an extremely difficult relationship with the US FDA and was the subject of a detailed and damning FDA (Enforcement) Warning Letter on April 8, 2014, and when they refused to respond to this Warning Letter, the FDA then broadcast a worldwide Safety Notice on April 29, 2014, the FDA version of the Death Penalty. This second letter effectively ended Shasta's ability to be a product design specifier and manufacturer, due to a total lack of regulatory adherence in the highly regulated medical device industry. It is confusing to consider what Shasta could have possibly been thinking. The company's acquisition of Genstrip (now GenUltimate!) was fortuitous in its timing given the finality and outcome of Shasta Technologies' fatal troubles with the FDA.

The worldwide market for at-home blood glucose testing is an estimated $17.2 billion, inclusive of the 2013 and 2016 changes to the Federal Medicare programs which gutted almost one-third of the U.S. market. The current GenUltimate! competes directly with one of the largest worldwide platform manufacturers the venerable Johnson & Johnson (Lifescan Inc.). GenUltimate! (and the earlier GenStrip 50) were developed for use with the OneTouch Ultra legacy system for at-home blood glucose testing, a

system currently used daily by over 3 million diabetes afflicted Americans and 5.8 million diabetics world-wide. GenUltimate! competes in the overall at-home testing market by offering an economical solution to former users of the legacy platform provider's product. The company's GenUltimate! product, designed to meet new European Union standards is a much improved version. Our business model is unique to this market channel as our major business focus is directed toward diabetics who have attempted a change of their glucose monitoring platforms (systems) or those currently using the J&J legacy products but are dealing with escalating prices and lower insurance reimbursements. At the time of the introduction of GenStrip in March 2013, J&J controlled just under 30% of this market and 100% of its own Lifescan, Inc. OneTouch Ultra market.

Throughout 2012 in anticipation of the introduction of Genstrip, we evaluated our brand-name distribution model, a model that provided streams of revenue but extremely low profit margins, and over the course of the last 30 months we phased out sales of those brand name products that had been a backbone of our distribution business. In addition the brand name products distribution business created a situation where we had been distributing legacy products that competed directly with our GenUltimate! Phasing out these brand name products lowered our order (revenues) intake but allowed us to become a manufacturer, at a higher level in the greater market channel.

The company will continue to direct its marketing efforts to ambulatory and semi-ambulatory older Americans afflicted with diabetes and complications caused by diabetes and old age. The company, originally a medical IT company with proprietary IT product lines, acquired its medical products distribution business in late 2004 through a merger with Phoenix, Arizona based CareGeneration, Inc. We have grown the original CareGeneration concept through subsequent acquisitions of private businesses and strategic partnerships with larger private pharmacies.

We began our transition into these medical products channels on November 1, 2011 when we completed the acquisition of Diagnostic Newco LLC from its owner Kimberly Binder. Diagnostic Newco LLC was a design company that specialized in product packaging design, medical products advertising design and graphic art. Ms. Binder has joined the staff of the company's Pharma Tech Solutions, Inc. subsidiary specifically for these purposes, and has worked closely with the contract manufacturers for GenUltimate!, making subtle changes to packaging design and more recently integrating the new FDA UDI product identification data system, among other responsibilities. She is also responsible for the package design for new diagnostic products the company is currently working on, including the GenSure! and the upcoming GenChoice! And GenPrecis! products. Ms. Binder is also owner of Genstrip Direct LLC and Full Circle Diabetes LLC, her own distribution companies, which she operates separately from her (Decision Diagnostics Corp. and Pharma Tech Solutions, Inc.) company related responsibilities.

We also intend to acquire additional private companies, or partner with small engineering companies that have developed technology requiring either regulatory approval, distribution expertise or both. In December 2011 we made another small acquisition, to acquire the services of Mr. Patrick Deparini. We are moving quickly to achieve our goal of becoming a vertically integrated, full service value added provider of products and services to an ever-growing market. The at-home diabetes testing market continues to grow as diabetics continue to be diagnosed and treated. The market for diabetes testing products is already in the tens of billions of dollars continues to grow rapidly. We also intend to make additional capital investment later in 2018 in our Korean contract manufacturer and advanced development partner for the manufacture of GenPrecis! and two new products GenAccord! And GenCambre!, both products that will compete with existing legacy products not previously mentioned.

The company's current proprietary product offering, cleared by the FDA for commercial distribution on November 30, 2012, and now in its later branded version, the GenUltimate! blood glucose diagnostic test strip for at-home testing. Genstrip, the original product, is a product originally conceived by Shasta Technologies LLC, who proved incapable of attaining the necessary regulatory approvals after two attempts, 2009 and 2010/2011. In addition the original Shasta concept could not clear the FDA 510K process on the basis of performance, and had to undergo major design changes and a new 510K application that was eventually sponsored by us. The original Shasta product was acquired by our Pharma Tech subsidiary on March 20, 2014, and fits into a diagnostic product niche, fitting nicely into the world-wide self-test (home test) market that has been growing at a 15% annual rate. Since GenUltimate! is a rather unique product offering, employing a brand name razor blade only model (diagnostic test strip) into a razor (diagnostic meter) -- razor blade (diagnostic test strip) market, the Genstrip 510(k) application made for unusual challenges for the FDA and an educational challenge and opportunity for the company. In fact, the company only recently (March 15, 2016) concluded its dealings with the FDA pre and post market review staff, an on-going process that was begun on a sour note by Shasta in October 2009. The company believes that upcoming product offerings such as the GenChoice! and GenPrecis! products, will also be regulated by the FDA but will go through a much smoother review and comment process, particularly since receipt of a directed landmark ruling by the U.S. FDA, covering our third party developed diagnostics (developed, in development and to be developed). Since the company plans additional similar products in the future for other diagnostic platforms, in fact a product announced still in the current reporting year, the Genstrip/GenUltimate! experience, however slow and unresponsive it was, has provided lessons and experience which is already being put to use.

Until our receipt of the landmark March 2016 ruling from the FDA, two years (and growing) was a standard development to market timeline for in-vitro diagnostic products similar to Genstrip / GenUltimate! In fact the long review periods and stifling performance standards established have contributed to a large decline in new products offerings in the industry since 2014. Nonetheless, we are confident that our new products will enjoy a much speedier FDA review process. As a result of previous delays and failures by Shasta Technologies in completing its FDA 510k approval application, and then problems Shasta encountered in prosecuting its two original applications with FDA staff, the company changed its contractual responsibilities and obligations in June 2011 to include program management, regulatory process management, management of the manufacturing forecasting and distribution processes, and new products planning and development. Further (eventually fatal) on-going problems encountered by Shasta, which on their face proved irresolvable, presented the company with an opportunity that we seized. On March 20, 2014 our Pharma Tech Solutions, Inc. subsidiary acquired the intellectual property, the marks, and the GenStrip cleared 510(k). Subsequently we accomplished a rebranding of the original Genstrip product (as GenUltimate!), built manufacturing protocols, implemented a robust Quality System throughout 2014 and 2015, and then developed the improved GenUltimate! product. GenUltimate! has become the only version of the original Genstrip line that will be packaged to conform with the FDA UDI standards, and was released as UDI compliant as of September 24, 2016. Manufacturing of Genstrip 50 ended and on-going sales continued under the GenUltimate! brand, and includes the FDA UDI packaging.

In June 2010 the company was approached by the largest retailer in the world for the distribution and sale of the Genstrip product, then about to enter the 510k regulatory review process, at over 5,000 retail stores worldwide. A contract with this retailer was negotiated in September 2010 and subsequently renegotiated and renewed in April 2011, and as soon as the retail contract was agreed to and as a means to conduct market research, the company began seeking pre-conditioned letters of intent (pre-orders) for Genstrip, while continuing the prosecution of the 510(k) application on behalf of Shasta Technologies before the FDA. Discussions with this retailer and other similarly situated retailers had been on a litigation induced hiatus since our litigation with Lifescan, Inc. began in earnest in late March 2013. Lifescan Inc., the diabetes testing division of Johnson & Johnson sued the company in three separate suits, all in Federal court, beginning in September 2011. These suits proved costly in that their intended purpose was to keep the Genstrip product off of retail market shelves. Until these suits were settled in May 2016, the company's marketing abilities were severely limited. The company believes there will be additional limitations as long as Johnson & Johnson spends large sums to discredit the company and its products. However, it should ne noted that Johnson & Johnson announced in January 2018 that their entire diabetic business (three divisions, multiple products) had been put up for sale, and offers for some or all of their businesses had been received. The sale of their Lifescan business, if it occurs, should change the diabetic testing field to a great degree, and should bring more positive views of our company in an industry that we have been competing in for six years.

The settlements we did achieve with J&J provided a hard-fought victory for the company, particularly since in 2015 Shasta had admitted to patent infringements of all three J&J diabetic medical device patents that were being adjudicated. We settled these lawsuits in a novel manner, where Johnson & Johnson paid the company a settlement amount, for lawsuits where the company was a defendant, a rarity in matters where the payor had initiated the strike suit in the first place. J&J, as a part of the settlement, also granted the company licenses to three J&J patents (including one patent that J&J subsequently lost through final action by the US Supreme Court), the larger value gained from this 5-year legal battle. In March 2016, prior to its settlement, the company's Pharma Tech Solutions, Inc. and Decision IT Corp. subsidiaries brought suit against Lifescan, Inc. in Nevada Federal court for patent infringement, the company alleging that Lifescan, Inc.'s OneTouch Ultra product was and had been infringing both of the company's patents. In March 2017, after a protracted battle with J&J where they tried to invalidate the company's lawsuit, the court in a major ruling agreed that the company will be allowed to move forward (a major victory so early in the suit) and will also be allowed to allege the Doctrine of Equivalents, a legal doctrine that would preclude J&J from twisting words through its pleadings and expert reports to escape justice. In April 2016 the company amended its original suit to include allegations under the Doctrine of Equivalents.

"The doctrine of equivalents is a legal rule in many (but not all) of the world's patent systems that allows a court to hold a party liable for patent infringement even though the infringing device or process does not fall within the literal scope of a patent claim, but nevertheless is equivalent to the claimed invention(s)."

Further, in January 2016 the US Supreme Court ruled that the Doctrine of Laches, a defense used by many Defendants in patent infringement suits could no longer be used. This ruling further deprived J&J of one of its most important defenses against the company's current patent infringement claims.

The Current Business

Currently the diabetes testing market is dominated by four large pharmaceutical manufacturers who provide very similar and equally focused products, selling at essentially equal prices. Genstrip's original introduction, even with the fits and starts, employed a business model different than those models employed by the major market players. Recent successes in the on-line marketplace has allowed the company to alter the market dynamics, lowering average price (which has occurred) or allowing for

increased testing by diabetics for a lesser price, thereby affecting all market segments. The company's major current market focus is to pharmacy chains, grocery chains with in-store pharmacies, large all purpose retailers with in-store pharmacies, and group buying and chain pharmacy organizations. Although this has been part of the company's plans in the recent past, the difficult litigation with Johnson & Johnson as well as the advent of the July 2013 and July 2016 changes to Medicare reimbursement (and followed by private insurers) and the October 2016 reimbursement engineering, pharmacy business models are now blurred. Thus the company successfully added on-line sales to its business model.

The company has also implemented a very successful "direct to diabetic" business model and has (independently or along with our distributors) executed on-line agreements with several of the largest retail chains, diabetic supply co-operatives, group purchasing organizations, as well as on-line mass merchandisers such as Amacon.com, Ebay, Walmart, Sears, Jet.com and approximately 900 other on-line cooperatives and product aggregators. The company considers this rapid adoption to be a huge success gained in a very short period of time.

In June 2017 we were notified by Amazon.com, the largest retail portal for our products where we sell approximately 20,000 boxes of GenUltimate per month, that the listings for our products had been "hacked" by ghost sellers -- individuals and people who listed our products, accepted orders and cash money from diabetics, but were unknown to the company. Oftentimes product was never delivered to the diabetic even after receipt of payment. This practice called freeloading (by Amazon) is not rare, but once started it is difficult to eradicate. The company had to replace almost 5,000 units of GenUltimate as a result of the freeloading. Further, while freeloaders had a cost basis of zero, legitimate sellers and distributors were forced to compete with these zero cost sellers. Prices for GenUltimate plummeted and by October 2017 the product in its largest portal declined on average by 35%.

With the assistance of Amazon, who themselves became a distributor of GenUltimate!, the company was able to overcome these issues. With the assistance of Amazon we reordered our selling practice, implementing base (floor) pricing and implementing real-time policing of listings. As a result, by mid-December 2017 we were able to overcome this freeloading practice. Prices have recovered about one third of the Fall 2017 decline. The company also eliminated many small distributors of GenUltimate! from the Amazon portal. While these actions had the effect of lowering sales in 1Q 2018, our margins and our sales levels are recovering.

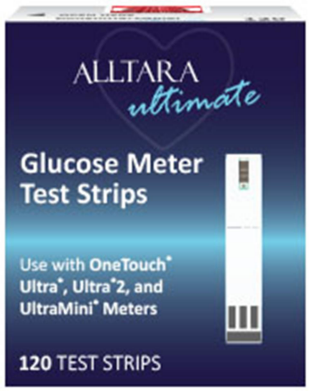
In March 2017 the company was contacted by the retail giant Walmart, who along with their acquired on-line retailer Jet.com, are attempting to duplicate and surpass the Amazon portal. Our GenUltimate! products have been sold on Walmart's (and Jet's) portals since November 2016. In the recent discussions Walmart offered us preferential listings on portals and Walmart Depot stocking at their regional transit facilities. GenUltimate! will now be sold by Walmart and shipments to diabetics will be fulfilled by Walmart. We accepted the offer (who wouldn't) and are changing our distribution agreement with Walmart (and Jet) in order that Walmart will sell and fulfill our products directly. Walmart customers who receive standing orders for their J&J Lifescan test strips will be a part of this new program. The company believes this to be a market enhancing deal since Walmart will become both a "push" and a "pull" retailer. No special pricing of our GenUltimate! pricing is required to implement this plan, owing, no doubt, to the footprint we have established on the on-line portals.

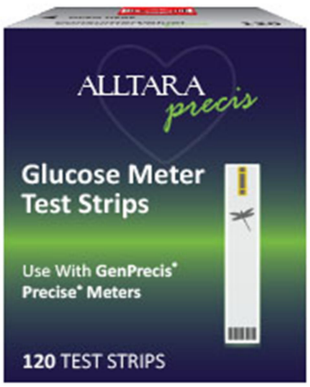
The company in the past has also offered information technology solutions in several medical care market channels by providing physicians with information at the point of care. Our products, unlike those from many other medical information companies, make use of smart cell phones such as the Apple iPhone, the Palm Pre, the Google Droid and a wide selection of Microsoft Windows based smart phones and operate in either in a wireless or "wired" mode, which allow physicians to carry, access and update their patients' histories, also known as electronic medical records or EMR, medication data, and best care guidelines - *all at the point of care*, or from any other location the physician may be located. In addition, the company's products employ proprietary mathematical game theory features adapted by the company for medical use that allow acceptance of diagnoses and treatment protocols where the medical information may have originated from one or several locations and one time or several times. Since the advent of "Obamacare," promising products like our own struggled to gain market acceptance in a reimbursement challenged market. The company cannot yet venture opinions or forecasts for its IT products now that the new administration While we have kept up with the evolving regulatory changes, we do not foresee implementation of our products and networks in the near future.

In October 2014 we adopted a value added/private label business model to address the issues brought to our market by the radical reimbursement changes by the federal Medicare program. We also hired a market executive with over 40 years of experience to implement our new strategy. We have doubled down on this strategy and now employ not only the services of the aforementioned expert, but also several of his partners and colleagues including the professional who put together the industry's "big box" pharmacy private label plan for diabetic test strips in 2006.

In March 2016 we also retained a product source company called Retail Monster, to represent our products to large drug chains ("big box pharmacy"), large retailers, chain grocers and the like. Unfortunately the arrangement with Retail Monster did not succeed, primarily because a group of company shareholders and persons claiming to be shareholders poisoned our relationship Retail

Monster by advocating, during repested calls, a "palace coup." After these incursions by shareholders and persons claiming to be shareholders, our relationship with Retail Monster remained cordial but destined to fail. The two companies decided to end the engagement on December 31, 2016. The efforts being expended in the "big-box" arena are greatly aided by the company's recent success with the explosively growing on-line Marketplaces, many sponsored by the large retail pharmacies and retail stores. These Marketplaces are fast growing sister organizations to these retailers, and typically not a part of legacy manufacturers marketing plans. The company's recent successes in the on-line Marketplaces has given the company a beachhead in this market as the uncertainty brought on by the J&J lawsuits has (finally) waned. In mid-March 2016 the largest US retailer agreed to raise the company's standing to the highest retail "rung" by offering a new supplier contract and in mid-March 2018 this retailer and its recently acquired wholesale products partner contacted the company and want to discuss a much enhanced relationship beginning in early April 2018.





Alltara *precis* is not yet available for sale or distribution in the United States or Puerto Rico.

Since March 2015 when we first we acquired special intellectual property and specialty manufacturing equipment which will shall serve our business interests now and into the future. We have increasingly turned to Alpha Capital Anstalt ("Alpha"), Navesink Device Initiatives, Sovereign Partners and Licgo Partners, whereby these organizations either purchased an 18-month 15% OID derivative instruments or Preferred C stock units, to facilitate the acquisition of intellectual property or manufacturing equipment, or to finance our growth. In 1Q, 2Q and 4Q 2016 and 2Q, 3Q and 4Q 2017 we completed additional financing transactions with both Alpha, Sovereign, and Licgo. Our most recent transactions with Alpha also financed an inventory credit line for the company so that we can meet many of the requirements of the largest retailers and maintain at least $300,000 in stock on hand at any time. Alpha also financed our acquisition of new specialty manufacturing equipment to facilitate our contract manufacturer in Korea as they develop our new GenChoice! product. The company will again turn to Alpha in April as we finance the completion of our GenPrecis! product.

In the Fall of 2014 the company announced its Discretion cloud wireless glucose monitoring product concepts, which will be manufactured for the company according to spec by its Korean contract manufacturer. In April 2015 the company entered into discussions with HMD Biomedical, Inc. in Taiwan for the importing of HMD's FDA cleared "Cloudia," product as a placeholder until the company's Discretion Messenger product for children would be ready. We ended our discussions with HMD Biomedical in October 2016, after determining that the "Cloudia" product was not robustly developed enough for North American markets and to further develop this product would require another 510(k) approval from the U.S. FDA which we did not wish to undertake. HMD Biomedical has not subsequently enhanced its Cloudia product, but the company has added many if not all of the creative features resident in the Cloudia to its GenPrecis! system.

The company has recently completed a further development of it's MD@Hand product, allowing diabetic users of the company's Discretion products to monitor and track their diabetes treatment and testing on their smart cell phones. The company plans at some point to spin-off its other MD@Hand and Residenceware technologies in a larger M&A transaction now in process.

The company entered into two international agreements throughout 2017. The first agreement, executed through the company's exclusive Korean agent, allows for delivery of the GenUltimate! and GenSure! (and certainly the GenPrecis! product when available) in quantity for sale in the Korean, Taiwan, Hong Kong, Vietnam and Thailand markets market. As of this writing, the Korean partners have ordered and paid for over 180,000 pieces (units/boxes) of GenUltimate! The company's second international agreement is through a South American financier who has businesses in Bolivia and Spain. This group initially placed a single two-year (term) order for approximately $17 million in GenUltimate! test strips, GenUltimate! meters and the company's new (2017) Firefly! Lancets. The South American financier also notified the company that he and those closely associated with him wished to subscribe to a $3.25 million to $5.0 million capital investment in the company. The group then signed and executed a Subscription Agreement for the company's Preferred D shares in April 2017.

After delivery of approximately 11,000 pieces (units) of GenUltimate!, 3,000 GenUltimate! meters and cases of lancets delivered to Bolivia, the company was contacted by authorities in the U.S. and then again several months later by regulators in Spain concerning the partners and silent partners involved with this international agreement. As a result of these contacts, the company, on March 20, 2017, terminated the Preferred D Subscription Agreement and terminated the International Distribution Agreement.

We have received multiple inquiries from companies interested in perhaps collaborating with the company for the implementation of its cell phone centric technologies MD@Hand and MD@Work. However, the market available for products similar to MD@Hand and MD@Work has changed since its introduction in 2009. The legal challenges to the new health care law and the federal government's inability to enact regulations have altered the landscape, again. We remain in discussions with multiple concerns for the marketing of our MD@ products, and any agreement we may enter will require us to provide contract software programming, providing a new source of revenue for the company. In addition to any proposed partnerships, we continue to discuss alternative propositions with other interested companies ranging from clinical laboratories, service organizations owned or aligned with medical health insurers, a medical content provider and legacy healthcare systems companies. There remains sustained interest in our MD@ technology. We may or may not entertain additional proposed partnerships for our implementation of the cell phone centric technologies, which has been hindered, as has the overall market, by the slow implementation of regulations, protocols and data formats by the Federal government, as well as a change in previously announced Federal government monetary incentives.

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Additional Background and Foundation

In May 2010, we entered into agreement with Shasta Technologies, Inc. and Broadtree, Inc. This agreement granted our Pharma Tech Solutions, Inc. subsidiary the exclusive marketing rights to a new diagnostic product not yet on the market named Shasta Genstrip ("Genstrip"). The Genstrip product was developed to compete against the market leader in the then $6.5 billion at home testing market. Shasta was in default of this 2010 Agreement within 90 days of its initiation. Penalties under that agreement and monies owed totaled in excess of $2 million in "delay" penalties, which they were unable to pay. In April 2011, the company renegotiated its agreement changing its many roles and adding responsibility for regulatory approval, manufacturing and forecasting, international sales and additional sales markets in the U.S. Shasta defaulted under this agreement as well. On March 20, 2014 we acquired the GenStrip intellectual property, its marks and the cleared 510(k). Shasta defaulted on this agreement as well. In addition Shasta breached or defaulted on two insurance settlement agreements, owing to the aforementioned J&J litigation. And finally, Shasta confessed to patent infringement of J&J's three patents.

On April 30, 2014 we first implemented our FDA mandated Quality Plan and are now operating as the manufacturer (operator) of the GenUltimate! test strip. We have implemented subsequent Quality Plans with our Korean contract manufacturer for our GenUltimate! product. Similar Quality Plans and FDA registrations will be in place for the company's GenChoice! and GenPrecis products in the near term, and for our GenAccord and GenCambre products later in 2018.

In August 2016 the company settled an insurance matter with Gotham Insurance, an IP Defense insurer, and Shasta covering legal fees associated with the 2011 and 2012 lawsuits brought by Lifescan, Inc. This settlement included a stipulation by Shasta to cease contacting and sharing confidential documents with persons who identified themselves as DECN shareholders. Several of these persons who contacted Shasta also contacted the aforementioned Retail Monster management. The stipulation gained in insurance settlement with Shasta does not preclude the company from pursuing Shasta, its principals and these "shareholders" in its omnibus lawsuit brought against Shasta et al. in 2014. The company is ammending its 2014 complaint to name additional Defendants including those persons who owned stock in the company who may have traded stock in the market based on information and documents provided by Shasta, or who were given confidential documents by Shasta, gained through the litigation discovery and provided to these shareholders, who then posted the information on public message boards.

We currently employ nine professionals at or locally managed through our executive business office located at 2660 Townsgate Road, Suite 300, Westlake Village, California 91361. In addition, we maintain two full-time and seven part-time positions located throughout the United States. We also maintain a Quality Assurance office through our exclusive agent in Seoul, Korea as a means to fulfill our quality commitments to the FDA. Our telephone number is (805) 446-1973 and our website addresses are and www.pharmatechsolutionsinc.com and www.genultimate.com. and www.decisiondiagnostics.com. Additional web sites will be added for our GenChoice! product (site in development) and our GenPrecis! product in the Spring of 2018.

As a part of the company's strategic plans, we have applied (to register) for twelve Trademarks with the USPTO. The company's Genstrip product is a registered Trademark of Shasta Technologies LLC. Our applications were filed with the USPTO in 1Q and 2Q 2015 and throughout 2016 and 2017. The company intends to use these Marks, as granted, to brand new products,

rebranding of existing products, and the establishment of a family of Marks associated with our company and its place in our industry. As December 31, 2017 the company has received registration confirmation from the USPTO for the following Marks:

"Alltara!"
"GenUltimate!"
"GenSure!"
"GenChoice!"
"GenAccord!"
"GenCambre!"
"GenPrecis!"
"Firefly!"
"ConsumerValue!"
"Infatig"
"Medicius!"

"Full Spread Electrode Technology"

Our marks for Alltara!, ConsumerValue!, Infatig!, and Medicius! will be used for product families as an integral part of our relationships with the "big-box" entities.

Beginning in the 4th Quarter 2015 and through 2nd Quarter 2016 the company suffered severe inventory shortage of the Genstrip 50 product at various times, owing to the timing of the various settlements with Johnson & Johnson by Shasta and a contract manufacturer, Conductive Technologies, Inc. For some period of time Conductive was unable, due to their settlement with Johnson & Johnson, to ship to the company certain quantities of the Genstrip 50 product. This problem began to clear up in late May 2016, and with the advent of adding the GenUltimate! product from Korea, shortages have been alleviated. The company's capacity for GenUltimate! production is now 625,000 packages per month (50 count and 100 count packages), for the new GenSure! product 250,000 packages per month (25 count and 50 count packages) and the new GenChoice! product (initial) 150,000 packages per month (50 count and 100 count packages). Recently, a mega-retailer has requested minimum inventories of finished product of 350,000 units/boxes. We expect other retailers to make similar requests. The manufacture of GenUltimate! and GenSure! are very similar and this capacity can be viewed as interchangeable. Similarly the manufacture of GenChoice! will be similar to the manufacture of GenAccord! and GenCambre!

The company's stock currently trades on the OTCMarkets OTC Pink Current tier of the market. The company's shares are DTC eligible. On May 12, 2015 the company made an application for a tier change to the OTCQX (common) tier. When the company's common stock fell in price beneath the $.10 threshold, and when our sponsoring broker shuttered his operation, our application went into hiatus. Subsequently, the company received direct communication from OTCMarkets concerning a new uplist program offered, beginning May 18, 2017, whereby the company might uplist within the OTCMarkets tiers as a Current Alternative reporting company and filer.

Business activities throughout the next twelve months:

The company's business on a day-to-day basis includes the distribution of our GenUltimate! products, (50 count and 100 count versions), distribution of our GenSure! product (25 count and 50 count versions) and later in 2018 our GenChoice! (25 count, 50 count and 100 count versions), and the GenPrecis! (25 count, 50 count, 100 count versions and a meter). Also within 120 days of this writing, the company will introduce its GenChoice! product which has recently concluded its clinical analyses. Our GenSure! will be sold only in certain International markets. In the next 120 days the company will have concluded the clinical analyses and filed for 510K clearance for its GenChoice! product (25 count, 50 count and 100 count versions). The GenChoice! product will be sold worldwide. Within 180 days of this writing the company will have concluded the clinical analyses and filed for 510K clearance for its GenPrecis! product (25 count, 50 count and 100 count versions and a meter designed with young diabetics in mind). The GenPrecis! product will be sold worldwide and will, most likely, require a strategic partner.

In mid-2017 the company embarked on an ambitious plan to re-brand all of its products, existing and upcoming, to sell into what is more commonly known as the private label marketplace, or the co-brand markets. These markets overlap to a high degree with what is also historically known as the "big-box" market. The rebranding contingency eventually grew to change the entire scope of our products developed for private label sales. In traditional diabetic supplies markets the packages had to include claims made in the original 510K application, plus new international symbiology and UDI identification. Packaging of the products was typically

designed to accommodate the capacities of the automation that packaged the products themselves. There was no magic involved with packaging. The 25, 50 and 100 count packages sold by the entire industry grew out of the capabilities of the automated packaging machines, not some grand plan. The entire industry became "me-too." The insurance reimbursement models associated with these 25, 50 and 100 count packages (overwhelmingly 50 count boxes) arose for the same reasons.

Companies in the manufacturing and marketing channels in the industry all employ these packaging processes, including Korean, Chinese and Taiwanese manufacturers. In truth, the manufacturer operations collectively decided not to pay an extra $10,000 for the packaging machines, or the $.10 for a slightly larger strip vial. The company believes this "me-tooism" to be a form of mental blinders. In implementing the company's new private label strategy, Decision Diagnostics decided not to bow to the packaging machine or "me-too" limitations. Instead the new packaging to be employed by the company will take into account diabetic testing patterns and the average number of testing days in a month. Private label versions of the company's products will be packaged in sizes of 30 count, 60 count and 120 count packages. This concept has been readily accepted by the company's private label target list in a detailed survey, and it is believed that this new packaging concept will be a marketing coup. Sales to the private label industry will be through private label product groups where every private label partner will own a private label group, each group containing all of the company's products in selective private label packaging.

The company currently has three major private label targets, the largest drug store chain, another top-5 drug store chain, and the second largest grocery store chain. In addition, the private label packaging is being offered to the largest drug store chains in Mexico and Canada. The Mexican chain, who also has numerous stores in Chile and Argentina has moved quickly. However, in all cases the sales process is in the closing stages. Closes of this nature, do however, take time. The company has Trademarked four product label groups for exclusive sales of products to the private label concerns: Alltara!, Advant!, Infatig!, and Medicius!.

Beginning in November 2009, we introduced our cell-phone centric medical IT products that offer solutions in medical care and management by providing physicians with information at the point of care. Unlike other medical information systems using standard computer terminals or even palm-sized computers (PDA's), our software applications operate on a series of late generation smart e-cell phones including the Apple iPhone, the Palm Pre, the Google Droid, several makes of RIM's Blackberry and many versions of the Microsoft Windows smart phones. Our products allow physicians to access and update their patients' histories, medication data, and best care guidelines - *all at the point of care*. The company's Electronic Medical Records software is believed to be the first EMR application running on any palm sized mobile device. Recently we ported our software to run on a series of pad computers such as Apple iPad and the 'Droid powered pads. We eagerly await the new version of the national health plan, which might finally create markets for our products.

Our 12-month business objectives include:

1. The practice of specializing in the distribution of GenUltimate! and GenSure! products, and the completion of the GenChoice! and GenPrecis! products. We also intend to add several brand-name medical diagnostic and medical disposable products (lancets through our Firefly! Product, as well as several lines of insulin syringes and pen needles, all associated with the on-going care of diabetes-inflicted patients, and the world-wide distribution of our proprietary diagnostic products.

2. Combining our wholesale and retail diagnostics distribution with the major successes we have had in the on-line retail markets, and adding legacy retail organizations (already some legacy retailers of note). See discussion above concerning private label opportunities and our private label lines.

3. Continue to implement the plans provided by our agent MWK LLC, and secure big-box pharmacy chains, chain grocers and nationwide retailers in addition to the private label groups previously discussed.

Recent Business Milestones:

In 2017 the company has accomplished the following milestones.

1. We completed the design and manufacture of GenSure! glucose test strips for the international markets, and completed development of our GenChoice! and GenPrecis! products.
2. We began patient clinical and clinical trials of two new test strip products, our GenChoice! and GenPrecis! test strips and the GenPrecis! Precise meter.
3. We are pressing our suit against Johnson & Johnson and several divisions for manufacturing products that infringe on

our patents. We won a major early battle in this suit where the trial judge granted us the opportunity to argue the Doctrine of Equivalents, an important concession in this case given J&J's penchant for the twisting of words and drawing lines through random dots. This suit began its prosecution phase on March 15, 2017 with the trial judge's early ruling. We await an important follow up ruling that if in our favor could bring about an end to the case.

4. The company initiated a marketing program to the on-line Marketplaces sponsored by pharmacy chain, department store and grocery store retailers, as well as mass merchandisers, and including the largest retailers. This program has so far been the most successful endeavor since our inception.

Financing Requirements

At December 31, 2017, we had cash of $1,088,761 and negative working capital of $545,337. We anticipate that we will require $72 million in trade debt financing (trade debt is conventional debt, not equity or some equity instrument) to finance our expected sales of our *"Gen"* products, as the current litigation ended in the company's favor. In March 2012 we renewed our agreement with Alpha Credit Resources ("ACR") to obtain this debt financing. In November 2013 we executed a new line of credit with Alpha Credit Resources, replacing our pervious line. The new credit line was for $12.5 million, but with the velocity of our product sales, could have yielded over $250 million in annually available credit. We never drew down any credit financing from ACR. Subsequently ACR became a part of major legal problems and is in liquidation. The company has unsuccessfully tried to replace ACR with a similar instrument. We plan to interview several other companies in the coming months.

Additionally, in order to accelerate our growth rate and to finance general corporate activities, we may supplement our existing sources of funds with financing arrangements at the operating system level or through additional short-term borrowings. As a further capital resource, we may sell or lease certain rights or assets from our portfolio as appropriate opportunities become available. However, there can be no assurance that we will be able to obtain any additional financing, on acceptable terms or at all.

Liquidity and Capital Resources

A critical component of our operating plan impacting our continued existence is the ability to obtain additional capital through additional equity and/or debt financing. We do not anticipate generating sufficient positive internal operating cash flow until later in 2018, as a result of several factors, including our on-going litigation with a division of Johnson & Johnson, and the change in our status from exclusive distributor of the GenStrip, to the manufacturer of this and similarly situated products, complete additional financial service company acquisitions and generate substantial revenues, which may take the next few years to fully realize. We believe we are adequately capitalized in the near term, but as our *"Gen"* products grow along their product life cycles, we may not obtain the necessary capital to pursue our strategic plan, and in the ultimate negative situation, we may have to cease or significantly curtail our operations. This would materially impact our ability to continue operations.

As of December 31, 2017, we had cash and cash equivalents of $1,088,761, inventory of $316,659, prepaid expenses of $859,413, and accounts receivable of $437,904. Net cash used by operating activities for the twelve months ended December 31, 2017 was approximately $1,558,919. Current liabilities of $3,248,074 consisted of: $805,555 of accounts payable and accrued liabilities, accrued interest of $173,433, and notes payable of $2,029,087. As of December 31, 2017, we have a negative working capital of $545,337.

Contingencies and Litigation

We transact commerce in several medical products market channels. We also transact commerce by licensing our proprietary medical software that functions by moving confidential medical data through our proprietary medical information technology devices and networks. Our original Genstrip product required initial regulatory approval by the USFDA as well as on-going USFDA approvals during the product life cycle. Further, Genstrip required medical patient trials and competes directly with a major platform manufacturer. We insure against any claims made against the company for our Genstrip product.

Our GenSure product is sold only in international markets. We are protected against claims of patent and/or trademark infringement by virtue of our 2016 settlement agreement with Johnson & Johnson and two J&J divisions.

Our GenChoice! and GenPrecis! products will be sold worldwide. The company will have to protect against claims of infringement for both of these products. Patent and trademark infringement suits are often filed for strategic business reasons, having only a passing relationship to the patents or trademarks claimed to be at issue.

Healthcare, especially those segments where the company competes, is also very litigious. Competing companies often use litigation as a marketing tool, bringing litigation as a means to protect market share and limit market exposure. The medical industry is also intertwined. From time to time, we may become involved in claims and litigation that arise out of the normal course of business, such as litigation that emerges from disputes over damaged, missing or contaminated product, litigation that arises over payment disputes or claims of fair value. We may also become involved in disputes that arise over the business or business practices of our suppliers, payers and customers. It is not uncommon in our industry to find that a litigant has filed claims in multiple jurisdictions involving the same transaction or a single transaction. The company maintains substantial insurance coverage against suits that may arise over issues of damaged, recalled or counterfeit product and other product liability issues. The company has also been a victim of the unapproved acts of prior management. These acts have resulted in claims from individuals and entities since the Board relieved former management of duty in 2006. Nonetheless, these claims have resulted in the use of management time and company resources to investigate, litigate, or settle. In addition, the company accrues contingent legal fees and product liability fees. As of December 31, 2017, our accrual was $485,069 and $245,069.

From time to time, the company may also be subject to demands from individuals or entities. These demands and disputes may consume management time and company resources. Other than as noted below, if there is such a disclosure, there are no pending matters at the current time that in management's judgment may be considered potentially material to us.

We were in litigation with Lifescan Inc. a subsidiary of Johnson & Johnson beginning in September 2011. Lifescan had maintained throughout that our Genstrip (now known as GenUltimate!) product infringed on three of their patents. One of these patents became the subject of peripheral litigation activities, and two Appeals (one for each side) to the U.S. Appeals Court for the Federal Circuit (the patents appeals court). In January 2016 the Court of Appeals for the Federal Circuit ruled in its Mandate that this one foundational patent and the claims made by the assignee Lifescan, Inc. was struck (killed) due to obviousness (a clever wording meant to obscure a connection between the Lifescan, Inc. invention and earlier generation technologies dating back to the late 1970s). Throughout this Appeal process, and a litigation process waged through the USPTO, the company prevailed. In addition, as a result of certain claims and allegations made by Lifescan after the close of the USPTO final determination (in favor of the company), the office of the Solicitor General intervened against Lifescan Inc. in the Federal Circuit court and was of great assistance in getting the Lifescan, Inc. patent revoked. Nonetheless the seeming baseless allegations and claims made by Lifescan against the company have taken their toll, limited our ability to sell our GenStrip (now known as GenUltimate) to large entities ("big box stores") and greatly extended the court processes.

In the Spring of 2013, fearing the impact of the Genstrip product in an open market, Lifescan took it upon themselves to violate a court protective order and prepared and sent out thirty page certified (veiled threat) letters to customers of the company and the customers of the company's customers, making it clear to these entities that should they do business with the company, or buy Genstrip (now GenUltimate) product from others doing business with the company, they could or would be added as defendants to the patent infringement suit. Most independent pharmacies in the U.S. sell less than a case (24 boxes) of any single brand of glucose test strips monthly. It is easy to ascertain that an independent pharmacy would choose not to "poke the bear" and risk a several hundred thousand dollar defense, rather than halting sales of Genstrip. Some large retailers were visited or called by Lifescan management and provided with face to face veiled threats. Lifescan even calculated that by breaching the protective order, the sanctions they would be assessed would amount to far less than the business loss they would otherwise suffer. Slowly however, the litigation environment enjoyed by Lifescan changed.

In May 2016 the company became aware of a clause Lifescan had inserted in its Franchise agreements. This clause set a penalty structure whereby should any Franchisee also buy non-Lifescan products (but more clearly our GenUltimate) they would lose their access to product rebates, and in certain instances their Franchise. Once aware of these illegal tie-ins the company complained to the Federal government, and in January 2017, for the first time since the onset of litigation with J&J, the tie-in clause was globally lifted by J&J. During the pendency of the 2011 and 2012 lawsuits, Lifescan was guilty of a number of unethical practices. For example, in December 2014 counsel for Lifescan wrote a letter to the trial judge who was hearing all three patent matters. This letter outlined a series of issues involving Lifescan's lead damages "expert" during litigation proceedings. Lifescan's expert claimed educational and qualification credentials that were not true at the time of the "expert" testimony, and are not true even today. This expert also assisted Lifescan's counsel in at least one other case, and other companies' counsels in unrelated cases. Testimony from this expert, in each instance, allowed the Plaintiffs in these cases to secure court rulings to the detriment of the Defendants. In the company's case this expert was used twice and assisted Lifescan to receive preferential treatment from the court for setting of a litigation bond to cover potential damages, wherein the "expert" through testimony limited the scope and calculation of damages in the setting of the damages protection afforded by the litigation bond and the damages resulting from Lifescan's violation of the court protective order. Lifescan's letter admonition came over a year after their successful use of this "expert."

In March 2016 the company filed suit in the Federal District Court of Nevada against Lifescan, Inc., Lifescan Scotland, Ltd. and Johnson & Johnson, citing infringement of two patents owned by the company. After an exchange of demand letters and posturing by

the Defendants, including Defendant's Motion to Dismiss, the company prevailed in an important early determination by the trial judge. At a hearing in March 2017 the Federal judge denied Lifescan's Motion to Dismiss, granted the company's request to allege the Doctrine of Equivalents and set dates beginning in early April 2017 and ending in early November 2017 that could set the stage for a ruling. Sometime in June 2017 the company expects to amend its suit a second time and name other "infringers" as well as adding additional counts to the suit. Federal rules for patent infringement suits have changed, and these suits are now adjudicated over an 18-24 month period. The trial judge's ruling in mid-March seems to foot with this schedule. In addition, if the schedule set by the judge does not end the litigation, there are five scheduled Mediations in front of a Federal Judge Magistrate pushing the process along.

On May 20, 2016 the company settled all of Lifescan's patent infringement claims as well as the company's Anti-trust and false advertising counter-claims against Lifescan, Inc. and Johnson & Johnson. Neither side in these litigations was a clear winner. The company's products were artificially denied a market for almost 3 years, but on the other hand, the company did receive rare settlement monies and other compensation from Lifescan in a suit where Lifescan was the Plaintiff. The amount of the settlement monies received by the company was confidential, as is often the case when Plaintiffs dismiss or lose a complex case, but confidentiality aside, the entire settlement was structured as a license agreement whereby Lifescan, Inc. granted licenses to the company for its test strip patents in return for accommodations regarding the anti-trust and false advertising claims made by the company. The licenses to the Lifescan Inc. patents were of great value to the company in the overall settlement.

In March 2016 the company filed suit against Johnson & Johnson and two Lifescan divisions through our two IP subsidiaries. DECN filed the lawsuit in the United States District Court, District of Nevada, in Las Vegas, NV, Case 2:16-cv-00564, titled Pharma Tech Solutions, Inc. et al v. Lifescan, Inc. et al naming Johnson & Johnson and its divisions Lifescan, Inc. and Lifescan Scotland Ltd. for alleged infringement in relation to U.S. Patent numbers 6,153,069, an apparatus patent, and 6,413,411, a method patent. The suit seeks at least $400 million in damages.

Fearful that the allegations in the suit were spot on, Lifescan filed a Motion to Dismiss which was denied. J&J, consistent with their historic tactical pattern of litigation delay, then filed a Motion for Summary Judgment. Despite a low probability of success, and the absence of legal appeal option for these Motions, J&J has through its filing successfully delayed the legal process for thirteen months to date. The trial judge has also ruled that PharmaTech would be permitted to file an amended complaint which could include further detail concerning patent infringement under the Doctrine of Equivalents; a significant advantage which minimizes the companies' burden in infringement cases. Once this Motion activity is concluded the company believes that the legal pendulum once again reverts in the direction of our potent legal position, where it should remain for the remainder of the litigation.

A. Date and State (or Jurisdiction) of Incorporation:

INCORPORATED IN THE STATE OF NEVADA ON MARCH 2, 2001 AS ATR SEARCH CORPORATION

B. the issuer's primary and secondary SIC Codes;

5122, 7371

C. the issuer's fiscal year end date;

DECEMBER 31

D. principal products or services, and their markets;

GenUltimate! Glucose Test Strips for use with Johnson & Johnson Lifescan glucometer, GenSure!, GenChoice and GenPrecis! glucose test strips and meters. MD@Hand medical communication and EMR software for use with smart cell phones.

7) Describe the Issuer's Facilities

We currently maintain an executive office at 2660 Townsgate Road, Suite 300, Westlake Village, CA 91361. The space consists of approximately 2,300 square feet. The monthly rental for the space is $2,170 per month on a month-to-month basis. We also maintain a Quality Assurance office in at the facility of our exclusive manufacturer's representative in Seoul, Korea. We contract for space in a specialty public warehouse in Miami, FL, which serves as our importing, exporting and shipping and receiving terminal.

8) Officers, Directors, and Control Persons

The goal of this section is to provide an investor with a clear understanding of the identity of all the persons or entities that are involved in managing, controlling or advising the operations, business development and disclosure of the issuer, as well as the identity of any significant shareholders.

A. Names of Officers, Directors, and Control Persons. In responding to this item, please provide the names of each of the issuer's executive officers, directors, general partners and control persons (control persons are beneficial owners of more than five percent (5%) of any class of the issuer's equity securities), as of the date of this information statement.

Our executive officers, directors, and key employees are:

Name	Age	Position
Keith Berman	64	CEO, CFO and Director
Robert Jagunich	71	

Our shareholders elect our directors and our Board of Directors appoints our officers. As of the date of this filing, we have not held an annual meeting. All current directors have been held over until such time the annual meeting is held. Vacancies in our board are filled by the board itself. Set forth below are brief descriptions of the recent employment and business experience of our executive officers and directors.

Keith Berman has served as Chief Financial Officer, Secretary, Treasurer and Director of the Company since January of 2003. He was appointed CEO in July 2017. Mr. Berman has been involved in the development of in-vitro diagnostic products, dry chemistry products, and healthcare software including Intranet and Internet systems for the past 42 years. From July 1999 to present, Mr. Berman has held the position of President, founder and director of Caredecision.net, Inc. a private company engaged in e-health technology development. From March 2001 through June 2002 Mr. Berman also held the Position of President and Director or Medicius, Inc. From January 1996 to June 1999 Mr. Berman was the President and founder of Cymedix, the operating division of Medix Resources, Inc., now Ramp Corp. (RCO). Cymedix was a pioneer company in what was then known as i-health (Internet healthcare) now the e-health industry. Mr. Berman's professional background provides the Company with business management experience and an in depth knowledge of our industry. Mr. Berman received a BA in 1975 and an MBA in 1977, from Indiana University.

Robert Jagunich has served as a Director of the Company since January of 2003. Mr. Jagunich has 27 years of experience in the medical systems and device industry. From August 1992 to present, he has held the position of President at New Abilities Systems, a privately held manufacturer of advanced electronic systems used in rehabilitation. He also provides consulting services to companies such as Johnson and Johnson and has served as a senior executive in such publicly held companies as Laserscope and Acuson. From April 1996 to December 1997 Mr. Jagunich acted as a director of Cymedix Corporation, the operating entity of Medix Resources, Inc., and later, Ramp Corp. (formerly AMEX:RCO). Mr. Jagunich's professional focus on medical devices as well as the professional relationships he has developed throughout his career provides the Company with opportunities to expand current markets and utilize additional product resources not previously available. He received his BS in 1969, and his MS and MBA in 1971, from the University of Michigan.

Mr. Berman, officer and director, works full-time for the company. Mr. Jagunich attends meetings of the board of directors when held and provides 10% and 15% respectively of his business time in professional capacities to the Company.

The following table sets forth information the remuneration of our Principal Executive officer for the years ended December 31, 2016, 2015 and 2014:

Summary Compensation Table

Name and Principal Position	Year	Salary ($)	Bonus ($)	Stock Awards ($)	Option Awards ($)	Non-Equity Incentive Plan Compensation ($)	Nonqualified Deferred Compensation Earnings ($)	All Other Compensation ($)	Total ($)
Keith Berman,	2015	$ -0-	-0-	$ -0-	-0-	-0-	-0-	-0-	$ -0-
CFO and PEO	2016	$ -0-	-0-	$ -0-	-0-	-0-	-0-	-0-	$ -0-
	2017	$ -0-	-0-	$ -0-	-0-	-0-	-0-	-0-	$ -0-

Mr. Berman has served as Chief Financial Officer since January 2003 and as Principal Executive Officer since August 2006. During the fiscal years ended December 31, 2016 and 2015. Through September 30, 2017 Mr. Berman has received no cash compensation. Mr. Berman has not received any form of compensation as a result of our limited cash flow; Mr. Berman has agreed to accept stock awards as his compensation until such time the Company has the necessary resources available to provide a traditional compensation plan.

Name of Beneficial Owner, Officer or Director	Number of Shares	Percent of Outstanding Shares of Common Stock
Keith Berman, Chief Financial Officer and Director	480,103	<1.0%
Robert Jagunich, Director	929,301	<1.0%
Directors and Officers as a Group	1,409,404	1.28%
Barbara Asbell (founder) 7061 Los Coyotes Camarillo, CA 93012	1,162,590	1.05%
Directors, Officers and Beneficial Owners as a Group	2,571,994	2.34%

B. Legal/Disciplinary History. Please identify whether any of the foregoing persons have, in the last five years, been the subject of:

1. A conviction in a criminal proceeding or named as a defendant in a pending criminal proceeding (excluding traffic violations and other minor offenses);

   None

2. The entry of an order, judgment, or decree, not subsequently reversed, suspended or vacated, by a court of competent jurisdiction that permanently or temporarily enjoined, barred, suspended or otherwise limited such person's involvement in any type of business, securities, commodities, or banking activities;

   None

3. A finding or judgment by a court of competent jurisdiction (in a civil action), the Securities and Exchange Commission, the Commodity Futures Trading Commission, or a state securities regulator of a violation of federal or state securities or commodities law, which finding or judgment has not been reversed, suspended, or vacated; or

   None

4. The entry of an order by a self-regulatory organization that permanently or temporarily barred suspended or otherwise limited such person's involvement in any type of business or securities activities.

None

C. Beneficial Shareholders. Provide a list of the name, address and shareholdings or the percentage of shares owned by all persons beneficially owning more than ten percent (10%) of any class of the issuer's equity securities. If any of the beneficial shareholders are corporate shareholders, provide the name and address of the person(s) owning or controlling such corporate shareholders and the resident agents of the corporate shareholders.

None

9) Third Party Providers

Please provide the name, address, telephone number, and email address of each of the following outside providers that advise your company on matters relating to operations, business development and disclosure:

Administrative Legal Counsel
Name: Thomas C. Cook
Firm: Law Offices of Thomas C. Cook
Address 1: 8250 W. Charleston Blvd. Ste. 120
Address 2: Las Vegas, NV 89117
Phone: (702) 242-0099
Email: tccesq@aol.com

Error Repair

The company will endeavor to repair any and all errors that new sets of eyes find in this document after its posting, whether these errors are in spelling, grammatical, punctuational or numeric. We are not perfect.

10) Issuer Certification

The issuer shall include certifications by the chief executive officer and chief financial officer of the issuer (or any other persons with different titles, but having the same responsibilities).

CERTIFICATION

I, Keith Berman, certify that;

(1) I have reviewed this disclosure statement and Annual Reports for the periods ended December 31, 2017 and December 31, 2016;

(2) Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

(3) Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

(4) The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

(a) designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

(b) Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

(c) revaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

(d) disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

(5) I have disclosed, based on our most recent evaluation of the internal control over financial reporting, to the registrant's auditors and the audit committee of registrant's board of directors (or persons performing the equivalent functions):

(a) all significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

(b) any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: March 30, 2018

/s/Keith Berman
Keith Berman
Principal Executive Officer and a Director
(Principal Financial Officer)

CERTIFICATION OF PRINCIPAL EXECUTIVE OFFICER AND PRINCIPAL FINANCIAL OFFICER
PURSUANT TO 18 U.S.C. SECTION 1350,
AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

The undersigned, Keith Berman, the Principal Executive Officer of Decision Diagnostics Corp., and Principal Financial Officer of Decision Diagnostics Corp.,., hereby certifies, that, to his knowledge, the Annual Report of Decision Diagnostics Corp. for the periods ended December 31, 2017 and December 31, 2016, fully complies with the requirements of this Disclosure Statement and of Section 13(a) or 15(d) of the Securities Exchange Act of 1934, as amended, and that the information contained in the Annual Report and this disclosure fairly presents in all material respects the financial condition and results of operations of Decision Diagnostics Corp. and its subsidiaries.

Date: March 30, 2017

/s/Keith Berman
Keith Berman
Principal Executive Officer and
Principal Financial Officer

A signed original of this written statement required by Section 906, or other document authenticating, acknowledging, or otherwise adopting the signatures that appear in typed form within the electronic version of this written statement required by Section 906, has been provided to Decision Diagnostics Corp. and will be retained by Decision Diagnostics Corp. and furnished to any regulatory body or OTC Markets, Inc. or their staff upon request.