

International Journal of Medical Science and Advanced Clinical Research (IJMACR) Available Online at: www.ijmacr.com

Volume – 1, Issue – 1, January – February - 2018, Page No. 13 - 15

An Analysis with Vitamin-B-12 - A Cancer Vitamin

Sitaraman Shekhar<sup>1</sup>, Ramagopalan Shetty<sup>2</sup>, Madhuri Rav Reddy<sup>3</sup>

<sup>1,2,3</sup> Amala Institute of Medical Sciences, Amalanagar, Thrissur, Kerala, India

Correspondence Author: Madhuri Rav Reddy, Amala Institute of Medical Sciences, Amalanagar, Thrissur, Kerala, India

E-Mail id: madhurirr01@gmail.com

### **Conflicts of Interest:** Nil

# Abstract

Laetrile, semisynthetic form of amygdalin with a controversial history called by, cancer vitamin, vitamin-B17 that is widely promoted on the internet in many websites and social media as treatment for cancer grabbed my interest to search for possible scientific evidence and reports published on this vitamin-B17. Amygdalin with no proper clinical effective evidence and proofs on cyanide toxicity is until the day in popularity as anticancer vitamin and sold on line at expensive cost. Due to hydro cyanide group present in both amygdline and letrile they exhibit side effects as like cyanide poisoning.

**Keywords:** Vitamin,B-17, Amygdalin,Laetrile, US-FDA **What is amygdaline/laetrile** 

Amygdalin, first isolated in 1830 from bitter almond seeds (prunus dulcis) by Pierre-jean Robiquet and

Antoine Boutroun-charlard. Amygdalin is a cyanogenic glycoside found in plants belonging to family of Prunus. It naturally occurs in Prunus fruit, pips like apricot, bitter almonds, plum, raw nuts; it also is existed in clover, lima beans, and sorghum. Laetrile is a semi-synthetic form derived from Amygdalin. Laetrile (patented 1961) is an acronym derived from the terms laevorotatory and mandelonitrile<sup>[1-4]</sup>. Amygdalin/Laetrile

Both amygdalin and laetrile contain, mandelonitrile, which decomposes to benzaldehyde and hydrogen cyanide. Since 1950, several versions on developments of Laetrile as anti-cancer drug, reported and published. It was theorized in 1962 as "cancer protein" in a book, Krebs, Sr., The term vitamin B-17 was given to laetrile by Ernst Krebs Jr, he also said, cancer caused due to deficiency of vitamin-B-17<sup>[5-7]</sup>. In the 1950s, Ernst T Krebs in the U.S. developed an apparently non-toxic, semi-synthetic intravenous form of Amygdalin to treat cancer.

In 1845, amygdalin used as cancer treatment in Russia and reported positive results in the first patient treated<sup>8, 9.</sup> In 1970's laetrile gained importance as monotherapy for cancer and by 1978, in the United States, 70,000 people reportedly treated with Laetrile. With the buildup of many controversies around laetrile on ineffectiveness in cancer therapy and cyanide toxicity, US FDA has exempted it from approval.

### How does it acts as anticancer drug.....?

Laetrile (containing amygdalin or benzaldehyde), as cancer remedy is controversial until the day due to lack of sound clinical evidence. However, proponents claim that amygdalin is selectively effective against cancer cells by below mechanisms.

Some hypothetical theories say that, laetrile readily delivered to cancer cell along with glucose. Cancer cells rich in  $\beta$ -glucosidases enzyme, hydrolyses laetrile to release cyanide and benzaldehyde forming a killing targeted poison that attack neoplastic cells.

Corresponding Author: Madhuri Rav Reddy, Volume -1 Issue -1, Page No. 13 - 15

Normal healthy cells are unaffected as they contain less  $\beta$ glucosidases enzyme and high protective enzyme, Rhodanese (thiosulfate sulfurtransferase). Rhodanese found in healthy cells and lack in cancer cells, neutralizes hydrogen cynide and benzaldehyde in to utilizable compounds thiocyanate (rhodanide) and benzoic acid.

Some theories say that cyanide toxicity is due to gut microflora because, Laetrile on oral ingestion is metabolized by alkaline duodenal and intestinal juice enzymes forming D-glucuronic acid and L-mandelonitrile. L-mandelonitrile is further hydrolyzed to hydrogen cyanide and benzaldehyde thatcause cyanogenic toxicity. Hence, said that intra venous preparations of Laetrile are safe from cyanide toxicities<sup>10-11</sup>.

# Is it safe....?

On oral administration of laetrile/Amygdalin, intestinal microflora that releases cyanide from laetrile. Cyanide is a neurotoxin, causes serious adverse events, like, headache, dizziness, nausea and vomiting, dermatitis or, in severe cases, disturbed consciousness, tachycardia, respiratory distress, liver damage, coma and death. There are few evidence of case reports saying that intensity of cyanide toxicity increases with concomitant administration of vitamin-c, raw almonds, fruits or vegetables that contain beta-glucosidase (e.g., celery, peaches, bean sprouts, carrots).

Many cases reported on cyanide poisoning following ingestion of bitter almonds<sup>21</sup>. However, there are no proper evidences of cyanide toxicity on parenteral administration. Unfortunately, Laetrile is available as oral and intravenous, intramuscular preparations, and coming to treatment method, patients are treated initially for three weeks with intravenous or intramuscular injections followed by maintenance therapy with, oral administration of Laetrile<sup>[11-13].</sup>

# Clinical evidences of amygdaline for its anti-cancer activity

<sup>14</sup>In early 1970,'s a scientist named Dr. Kanematsu Sugiura, performed experimental tests on mice and claimed that laetrile effectively inhibited secondary tumors but has no effect on primary tumors. However, any scientist never reproduced the results after Dr. Kanematsu Sugiura.

### Conclusion

From the literature available from 1970's, there were no proven clinical effects on the use of Laetrile. In fact, all the studies highlighted cyanide toxicity. However, even today amygdalin is available in the market as oral preparations by naming as apricot seed powder. After many decades of clinical use of amygdalin in the field of alternative medicine to treat cancer, no sound clinical evidences were found on its anti-cancer activity.

#### References

[1]. Young JH. Laetrile in historical perspective. In Merkle GE, Petersen JC, editors. Politics, Science, and Cancer: The Laetrile Phenomenon. Boulder, CO: Westview Press, 1980.

[2]. Vetter 2000 Vetter J. Plant cyanogenic glycosides. Toxicon 2000; 38(1): 11–36.

[3]. Laetrile/Amygdalin. National Cancer Institute.

[4]. Lerner IJ. Laetrile: a lesson in cancer quackery. CA: A Cancer Journal for Clinicians. 1981. 31: 91–95.

[5]. Ellison NM, Byar DP, Newell GR: Special report on Laetrile: the NCI Laetrile Review. Results of the National Cancer Institute's retrospective Laetrile analysis. N Engl J Med 299 (10): 549-52, 1978.

[6]. Milazzo S, Ernst E, Lejeune S, Boehm K, HorneberM. Laetrile treatment for cancer. Cochrane Database ofSystematic Reviews 2011, Issue 11. Art. No.: CD005476.

[7]. The laetrile controversy. In: Moss RW: The Cancer Industry: The Classic Expose on the Cancer

# Madhuri Rav Reddy, et al. International Journal of Medical Sciences and Advanced Clinical Research (IJMACR)

Establishment. Brooklyn, NY: First Equinox Press, 1996, pp 131-52.

[8]. Cancer.gov. Complementary and Alternative Medicine for Health Professionals. United States: National Cancer Institute. March 15, 2017.

[9]. Rietjens, Ivonne M. C. M.; Martena, Martijn J.;
Boersma, Marelle G.; Spiegelenberg, Wim; Alink, Gerrit
M. (2005-02-01). Molecular Nutrition & Food
Research. 49 (2): 131–158.

[10]. Kennedy, Donald (1977). "Laetrile: The Commissioner's Decision" (PDF). Federal Register. Docket No. 77-22310.

[11].PDQ®Integrative,Alternative,andComplementary Therapies EditorialBoard.PDQLaetrile/Amygdalin.Bethesda,MD:National Cancer Institute.Updated <03/17/2017>.

[12]. Adele Stapf, Helen Cooke, Helen Seers, CAM-Cancer Consortium. Amygdalin/Laetrile [online document]. February 8, 2017.

[13]. Mohammed Helmy Faris Shalayel, Beyond Laetrile
(Vitamin B-17) Controversy-Antitumor Illusion or
Revolution, British Biomedical Bulletin ISSN 2347-5447,
2017, Vol.5 No.1:296.