

The viroses and virus-like diseases of the grapevine

A bibliographic report, 1971—1978

by

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Preview

This is the third of a series, "A bibliographic report" on progress in research and knowledge on the cause, nature and control of viroses and the virus-like diseases of grapevines. The first was "Bibliographie des Viroses de la Vigne, des origines à 1965", 1019 references to literature, by CAUDWELL, O.I.V. special publication 76 pp., 1965; and the second was "Les Viroses de la Vigne; Bibliographie de 1965 à 1970", references numbered 1020—1386, by CAUDWELL and text by HEWITT, BOVEY and CAUDWELL, *Vitis* 11, 303—324, 1972.

This report covers advances in knowledge expressed in literature from 1970 into 1978. It also has some omissions from prior dated literature. Efforts have been made to include as many as possible of the references in the literature on viruses, viroses, virus vectors, and virus-like diseases of grapevines. It is recognized that no doubt some references have been overlooked and that especially not all of the literature published in 1978 has been included.

This report contains references to papers presented at each of the last three meetings and published proceedings and/or abstracts of the International Council for the Study of Viruses and Virus Diseases of the Grapevine (ICVG), Colmar, 1970, Salice Terme, 1973 and Cordoba 1976.³⁾ The objective of the ICVG is to encourage on an international basis, research into the cause, nature and control of virus and virus-like diseases of grapevines. Researchers are encouraged to present results of research at international meetings of the ICVG, in seminars, formal and informal discussion sessions. Information concerning ICVG and the meetings may be obtained from the Secretary or any member of the steering committee. They are:

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³⁾ Although the Proceedings of this last meeting were published in 1979, they have been included in this bibliography. Some papers that are not included in the Proceedings are referred to as abstracts.

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The seventh meeting of the ICVG will be held in Ontario, Canada, in September 1980, H. F. DIAS is Chairman and may be contacted for information.

Introduction

The application of modern concepts into research on the cause, nature and control of grapevine and vineyard degeneration problems began only a relatively short time ago, 122, 124, 345, 347, 421, 482, 487, 488, 757, 964. (These numbers are references to literature in the "Bibliographie des Viroses de la Vigne des origines à 1965", OIV, 1965.)

During the years 1970 through 1978, great advances have been made in knowledge on the pathology of the degenerative virus and virus-like diseases of the grapevine. In reviewing the literature and progress it is interesting to note the activity of researchers, country by country, and the steps taken to analyze their local disease situation, review world knowledge, and proceed in research. The time lapse from the start of work as demonstrated in literature reviews, evaluations of vineyard problems, diagnosis of specific diseases and entry into research on development of new knowledge has in general decreased as the overall knowledge on the viroses and virus-like diseases has increased. Furthermore, progressive improvement in research on these grapevine and vineyard problems is evident in papers presented at the meetings of ICVG. Continuous and substantial improvement is demonstrated in reports on vine and vineyard performance, and quality of wine made from "healthy" and/or improved grapevine cultivars.

References

These references are of a more general nature, including reviews and commentaries, and some survey material.

Bibliographies: 1578 and 1993.

- 1388 — 1392 — 1396 — 1399 — 1405 — 1407 — 1410 — 1419 — 1422 — 1428
1431 — 1432 — 1446 — 1461 — 1474 — 1480 — 1497 — 1514 — 1517 — 1522
1528 — 1529 — 1532 — 1536 — 1538 — 1544 — 1545 — 1548 — 1549 — 1552
1553 — 1554 — 1555 — 1556 — 1559 — 1561 — 1563 — 1571 — 1591 — 1593
1626 — 1630 — 1638 — 1644 — 1658 — 1659 — 1669 — 1684 — 1685 — 1686
1693 — 1696 — 1709 — 1713 — 1715 — 1716 — 1725 — 1726 — 1727 — 1728
1730 — 1750 — 1752 — 1753 — 1760 — 1767 — 1784 — 1792 — 1794 — 1798
1804 — 1810 — 1814 — 1815 — 1816 — 1818 — 1819 — 1820 — 1851 — 1853
1856 — 1859 — 1860 — 1897 — 1902 — 1905 — 1906 — 1920 — 1921 — 1922
1940 — 1950 — 1954 — 1958 — 1959 — 1996 — 2009 — 2012 — 2015 — 2027
2030 — 2045 — 2046 — 2049 — 2052 — 2063 — 2079 — 2084 — 2085 — 2089
2091 — 2103 — 2104 — 2107 — 2109 — 2111 — 2112 — 2116 — 2117 — 2119
2123 — 2127 — 2129 — 2141 — 2143 — 2147 — 2154 — 2162.

Comments

It is necessary to retain the two groupings viroses and virus-like diseases within this general class of the diseases of grapevines. This is essential because viroses are caused by specific viruses that have been properly identified, whereas the virus-like diseases behave like and/or have the characteristics of a virus-caused disease but the pathogen of each has not been properly identified. In the past, some diseases were classed and referred to as virus diseases but the pathogens proved later to be other than viruses. For example, Flavescence dorée (FD) and Pierce's disease (PD) were in earlier literature termed or classed as virus diseases. It is known at this time that the pathogen of FD is a mycoplasma (1575, 1577, 1581), and that of PD to be a bacterium (1610, 1611, 1741, 1786, 2071). Perhaps in the reasonable future it will be possible to place each of the virus-like diseases under a pathogen class. Researchers should pursue in so far as practical the nature of the pathogens of the virus-like diseases. Knowledge of the pathogens of a disease opens the broader store of information and experience of the pathogen class to research on the grapevine disease.

Viruses and viroses of the grapevine

Following is a list of the viruses, viroses, and the reference numbers to literature in this bibliographic report. The literature shows that there are 25 viruses that have been isolated out of grapevines. Twenty-two of them have been identified and/or characterized and named, two had not been named and five as we know them appear to have been viruses of the grapevine for a long period of time; to wit, GBLV, GCMV, GFV, GLRV and possibly GJSV. Reference 1987 reports only that carnation ringspot virus was found in *Stellaria media* growing in vineyards with virus-like diseases. Viruses and viroses receiving the greatest attention are GFV, GLRV, TomBRV, TomRSV and AMV. There is good reason for high activity in research on GFV and GLRV — they appear to be distributed in vineyards over the world, and result in substantial losses.

References

Viruses isolated from grapevine and viroses. Common names of the diseases in grape in parentheses.

AlMV, Alfalfa mosaic virus

1510 — 1535 — 1537 — 1762 — 1890 — CMI/AAB 46⁴).

AMV, *Arabis* mosaic virus

1404 — 1505 — 1506 — 1509 — 1511 — 1512 — 1564 — 1605 — 1735 — 1762
1910 — 1939 — 1983 — 1997 — 2020 — 2050 — 2099 — 2151 — 2152 — CMI/
AAB 16.

AILV, Artichoke Italian latent virus

1762 — 1763 — 1820 — 1829 — 2002 — CMI/AAB 176 (= 1829).

BBWV, Broad bean wilt virus

1762

⁴) CMI/AAB Descriptions of Plant Viruses (with publication number). Edited by B. D. HARRISON and A. F. MURANT. Issued jointly by the Commonwealth Mycological Institute, Ferry Lane, Kew, Surrey, England and the Association of Applied Biologists.

Carnation ringspot virus (in *Stellaria media* in vineyards)

1987.

CM 112 virus

1645 — 1646 — 1838.

GBLV, Grapevine Bulgarian latent virus

1762 — 1820 — 1821 — 1822 — 1828 — 2090 — CMI/AAB 186 (= 1828).

GCMV, Grapevine chrome mosaic virus (Hungarian chrome mosaic)

1793 — 1805 — 1808 — 1824 — 1826 — 1827 — 1997 CMI/AAB 103 (= 1827).

GFV, Grapevine fanleaf virus (fanleaf, yellow mosaic and vein banding, court-noué, panachure)

1387 — 1391 — 1394 — 1400 — 1409 — 1416 — 1420 — 1421 — 1429 — 1430
 1434 — 1436 — 1437 — 1438 — 1439 — 1440 — 1441 — 1442 — 1452 — 1457
 1458 — 1495 — 1505 — 1506 — 1509 — 1511 — 1512 — 1513 — 1520 — 1521
 1557 — 1614 — 1622 — 1631 — 1632 — 1633 — 1638 — 1650 — 1651 — 1653
 1654 — 1655 — 1664 — 1667 — 1670 — 1674 — 1679 — 1681 — 1694 — 1695
 1707 — 1711 — 1712 — 1718 — 1724 — 1733 — 1735 — 1751 — 1759 — 1788
 1793 — 1801 — 1805 — 1808 — 1825 — 1835 — 1846 — 1849 — 1850 — 1852
 1854 — 1856 — 1858 — 1861 — 1865 — 1867 — 1878 — 1883 — 1898 — 1899
 1907 — 1909 — 1910 — 1911 — 1922 — 1926 — 1938 — 1939 — 1947 — 1948
 1950 — 1960 — 1964 — 1997 — 1999 — 2000 — 2003 — 2050 — 2052 — 2062
 2065 — 2066 — 2067 — 2068 — 2088 — 2100 — 2106 — 2108 — 2116 — 2126
 2138 — 2151 — 2152 — 2153 — 2154 — 2163 — CMI/AAB 28 (= 1733).

GFFV, Grapevine false fanleaf virus

1820.

GJSV, Grapevine Joannes-Seyve virus (Joannes-Seyve disease)

1617 — 1622.

GLRV, Grapevine leafroll virus (leafroll)

1407 — 1444 — 1445 — 1450 — 1515 — 1516 — 1520 — 1560 — 1625 — 1628
 1629 — 1638 — 1664 — 1680 — 1718 — 1751 — 1764 — 1788 — 1795 — 1800
 1801 — 1820 — 1825 — 1856 — 1867 — 1880 — 1883 — 1907 — 1922 — 1950
 1997 — 2023 — 2028 — 2041 — 2048 — 2052 — 2053 — 2054 — 2055 — 2056
 2057 — 2065 — 2096 — 2136 — 2161 — 2163.

PRMV, Peach rosette mosaic virus

1615 — 1616 — 1618 — 1623 — 1624 — 1942 — 1943 — 1944 — CMI/AAB
 150 (= 1618).

PotXV, Potato X virus

1691 — CMI/AAB 4.

PotYV, Potyvirus (Potato virus Y group)

1757 — 2053 — 2054 — 2055 — 2056.

RRV, Raspberry ringspot virus

1402 — 1506 — 1511 — 2039 — 2099 — 2151 — 2152 — CMI/AAB 6.

SMV, Sowbane mosaic virus

1923 — CMI/AAB 126.

SLRV, Strawberry latent ringspot virus

1509 — 1735 — CMI/AAB 126.

TMV, Tobacco mosaic virus

1474 — 1505 — 1509 — 1627 — 1758 — 1773 — 1856 — CMI/AAB 151.

TNV, Tobacco necrosis virus

CMI/AAB 14.

TRV, Tobacco ringspot virus

1687 — 1689 — 1802 — 1910 — 1998 — 2080 — 2083 — 2086 — CMI/AAB 17.

TomBRV, Tomato black ring virus

1402 — 1403 — 1505 — 1509 — 1512 — 1826 — 1835 — 1981 — 2021 — 2151
2152 — CMI/AAB 38.

TomBSV, Tomato bushy stunt virus

1762 — 1890 — CMI/AAB 69.

TomRV, Tomato ringspot virus

1459 — 1619 — 1620 — 1621 — 1688 — 1886 — 1998 — 2078 — 2080 — 2081
2083 — 2086 — 2087 — CMI/AAB 18.

GTomRV, Grapevine strain TomRV (Yellow vein)

1459 — 1696.

Viruses unknown or not identified

1856 — 2040.

Virus-like diseases of the grapevine

Following is a list of the virus-like diseases of grapevines and literature reference numbers. There are 17 listed, some may be synonyms. The truth will not be known until the pathogens are properly identified. Among these diseases, it appears that fleck (marbrure), legno riccio (stem pitting), corky bark, and yellow speckle are old diseases of the grapevine. They are latent in some cultivars and widely dispersed among the cultivars and rootstocks. Spread may have taken place as did that of GFV and GLRV, with nursery practices and the distribution of hybrid rootstock. Yellow speckle is latent in many cultivars when grown in California yet symptoms develop in the same cultivars when grown in parts of Australia. Legno riccio is latent in some cultivars and not in others, furthermore, there appears to be a scion/stock interaction for symptoms that show on some but not all combinations.

R e f e r e n c e s

Virus-like diseases: probably caused by viruses but not demonstrated to have a virus pathogen

Asteroid mosaic

1420

Bratislava mosaic

1922.

Corky bark (refer also to legno riccio, stem pitting, stem grooving)

1390 — 1525 — 1574 — 1666 — 1699 — 1722 — 1729 — 1734 — 1867 — 1883
1950 — 1952 — 2048 — 2094.

Chasselas latent (refer also to fleck)

1524.

Enation

1470 — 1690 — 1704 — 1706 — 1720 — 1801 — 1922 — 1929 — 1930 — 2066
2163.

Flat trunk

1729 .

Fleck (marbrure)

1400 — 1495 — 1632 — 1633 — 1718 — 1720 — 1724 — 1732 — 1790 — 1801
1847 — 1853 — 1856 — 1857 — 1867 — 1883 — 1894 — 1907 — 1922 — 1949
1951 — 1953 — 1960 — 1997 — 2096 — 2147 — 2163.

Grapevine summer mottle

1768.

Interveinal chlorosis

1856 — 1862.

Interveinal necrosis

1856.

Legno riccio (wood pitting, stem pitting, stem grooving); also refer to corky bark

1435 — 1443 — 1445 — 1447 — 1448 — 1449 — 1453 — 1454 — 1456 — 1520
1534 — 1636 — 1637 — 1638 — 1639 — 1640 — 1641 — 1666 — 1675 — 1676
1677 — 1699 — 1702 — 1705 — 1719 — 1729 — 1734 — 1791 — 1830 — 1835
1856 — 1900 — 1901 — 1922 — 2001 — 2003 — 2010 — 2051 — 2081 — 2083
2102 — 2113 — 2125.

Linear pattern

1856 — 1994.

Marbrure: see fleck

Rcaziteli yellow stipple

1451.

Shoot necrosis

1812.

Vein mosaic

1768 — 1790 — 1811 — 1856 — 1862 — 1915 — 1918 — 1951 — 1997 — 2147.

Vein necrosis

1790 — 1833 — 1853 — 1855 — 1856 — 1862 — 2147.

Yellow speckle

1451 — 1718 — 1866 — 2064 — 2065 — 2163.

Evidence would indicate that corky bark and legno riccio have the same pathogen because they each cause the same symptoms in the indicator plant LN33. However, this may or may not be so. Transmission tests from donors that induce common symptoms in a cultivar or even a group of cultivars (indicator plants) do not alone determine that the donors all had the same pathogen. For example, GFV, yellow mosaic strain and GCMV, and apparently a potyvirus also can induce common symptoms in several cultivars of grapevine. It will be necessary to determine the pathogen of some different corky bark and legno riccio diseased plants; then only if the pathogens are the same in each case are the names synonymous.

Complexes of viruses and/or pathogen combinations in grapevines

Classical examples of grapevine diseases composed of complexes of virus pathogens are infectious degeneration (*Dégénescence infectieuse*), Reisigkrankheit, and Roncet. Though GFV will be found most commonly in collections of the disease, there seems little doubt but that infectious degeneration in parts of each, France, Italy, Greece or elsewhere will have a different component of virus pathogens. The disease problems in any locality can be evaluated more accurately by proper diagnosis of causal viruses.

R e f e r e n c e s

1395 — 1397 — 1401 — 1406 — 1412 — 1413 — 1414 — 1421 — 1425 — 1426
1509 — 1564 — 1631 — 1681 — 2011 — 2086.

Surveys, general distribution and spread of the viroses and virus-like diseases

Surveys on the presence of grapevine virus disease problems in any country or locality are the first step in an analysis of vineyard health. These lead into research and vineyard improvement programs. Also after proper diagnosis, surveys establish distribution of the more specific diseases, set a basis for evaluating the health condition of vineyards, the potential for improvement, and support levels needed for research and amelioration programs.

Literature reports of surveys help in understanding world distribution of the viroses and potential for improvement in total production.

R e f e r e n c e s

The reader will also find survey literature in the references to the introduction, and later in the section on selection, sanitation and preservation.

1388 — 1397 — 1411 — 1421 — 1422 — 1461 — 1473 — 1511 — 1568 — 1648
1679 — 1681 — 1693 — 1749 — 1762 — 1792 — 1800 — 1897 — 1902 — 1907
1917 — 1919 — 1926 — 1995 — 1997 — 2015 — 2016 — 2017 — 2052 — 2079
2080 — 2103 — 2105 — 2111 — 2161 — 2162 — 2163.

Diagnosis: Indexing, transmission, trabeculae, and serology

Proper diagnosis is fundamental to any plant health program. The presence of signs such as vine degeneration, presence of internal signs like trabeculae and external symptoms are indicative of any disease condition. A syndrome in symptomatology gives a better diagnosis. Indexing by graft-transmission onto a set of indicator plants establishes transmissibility and virus-like nature of the causal pathogen(s). Sap inoculation to a class of indicator plants will further establish the potential virus pathogen. Electron microscopy provides information on morphology and more precise information on class and grouping of a pathogen. Serology will establish pathogen relationship and prove the presence of specific viruses and other pathogens. Final accurate diagnosis, however, requires isolation, purification, identification, reinoculation, establishment of disease and reputation as with the Koch's Postulate. Once an experienced researcher becomes familiar with a pathogen, its symptomatology in a host-range, etc., short cuts in diagnosis are justified.

R e f e r e n c e s

General

1393 — 1427 — 1433 — 1475 — 1496 — 1563 — 1565 — 1566 — 1567 — 1613
1652 — 1761 — 1916 — 1919 — 2095 — 2127 — 2139 — 2141 — 2144 — 2146.

Indexing by grafting on *Vitis* indicators

1574 — 1680 — 1699 — 1721 — 1866 — 1867 — 2024 — 2028 — 2037 — 2065
2088 — 2092 — 2112 — 2116 — 2130 — 2143 — 2146.

Mechanical (sap) transmission

1550 — 1841 — 1965 — 2031 — 2034 — 2053 — 2065 — 2088 — 2101 — 2133
2134 — 2135 — 2143 — 2152.

Serology

1506 — 1507 — 1508 — 1651 — 1695 — 1920 — 2035 — 2124 — 2140 — 2142
2143 — 2149 — 2150 — 2160.

Trabeculae

1504 — 1513 — 1653 — 1654 — 1655 — 1657.

Comparison of different methods

2065 — 2088.

Vectors of soil-borne viruses and nematodes

Viruses of some of the important and widely distributed viroses of grapevines are soil-borne and have nematode vectors. The importance and place of nematodes

as vectors of the soil-borne viruses is constantly enhanced in reports of research on the subject.

In the following the general papers on surveys and presence of nematodes in various countries and situations and soil treatments are listed. Included also are reports of research on specific nematodes and those found in vineyards with grapevine disease problems.

References

Nematodes, nematode vectors of soil-borne viruses, and soil treatments

General

1662 — 1774 — 1775 — 1776 — 1777 — 1782 — 1818 — 1957 — 2058 — 2059
2118.

Virus-vector relationships

1402 — 1404 — 1714 — 1802 — 1818 — 1957 — 2060 — 2061.

Nematode-surveys, soil treatments, fallowing

1408 — 1423 — 1463 — 1464 — 1465 — 1521 — 1546 — 1551 — 1562 — 1568
1592 — 1595 — 1599 — 1600 — 1601 — 1602 — 1603 — 1604 — 1606 — 1607
1608 — 1635 — 1679 — 1681 — 1682 — 1683 — 1711 — 1712 — 1738 — 1765
1766 — 1774 — 1775 — 1777 — 1778 — 1779 — 1780 — 1781 — 1783 — 1806
1808 — 1818 — 1836 — 1863 — 1864 — 1896 — 1931 — 1941 — 1945 — 1946
1948 — 1956 — 1966 — 1968 — 1969 — 1970 — 1973 — 1974 — 1975 — 1976
1979 — 1980 — 1982 — 1984 — 1985 — 1986 — 2018 — 2040 — 2043 — 2085
2115 — 2131 — 2137 — 2155 — 2156.

Xiphinema species

1417 — 1463 — 1464 — 1595 — 1735 — 1770 — 1864 — 1962 — 1966 — 1974
2001 — 2043 — 2156 — 2157.

X. americanum

1408 — 1703 — 1738 — 1802 — 1864 — 2066 — 2155.

X. diversicaudatum

1404 — 1564 — 1605 — 1778 — 1779 — 1780 — 1864 — 1983 — 2060 — 2099
2151 — 2152 — 2153 — 2154 — 2155.

X. index

1408 — 1425 — 1457 — 1458 — 1465 — 1519 — 1596 — 1598 — 1650 — 1703
1710 — 1711 — 1712 — 1738 — 1778 — 1779 — 1781 — 1783 — 1805 — 1808
1835 — 1836 — 1864 — 1924 — 1925 — 1928 — 1947 — 1948 — 1988 — 1990
1991 — 2018 — 2042 — 2060 — 2066 — 2068 — 2072 — 2126 — 2137 — 2151
2152 — 2153 — 2154 — 2155 — 2158.

X. italiae

1465 — 1778 — 1779 — 1780 — 1796 — 1864 — 2153 — 2154 — 2155.

X. mediterraneum

1778 — 1779 — 1780 — 1962 — 2072.

X. vuittenezi

1805 — 1807 — 1808 — 1864 — 1967 — 1977 — 1978.

Longidorus species

1402 — 1403 — 1463 — 1770 — 1778 — 1779 — 1780 — 1981 — 2099 — 2155
2157.

Trichodorus species

1770.

It is interesting to note that investigators continue to report *X. vuittenezi* associated with grapevine degeneration problems, but that no reports demonstrate the nematode to be a vector of one of the viruses of grapevines. Reports demonstrate that vector-nematodes may acquire and may transmit virus in a short period of feeding, hold and transmit virus over long periods if not for the entire adult life. Sites of virus retention by *Xiphinema* spp. appear in that portion of the alimentary tract shed in molting, and/or retained in adults.

Control of nematodes by chemical soil treatment is accomplished, but not without problems apparently related to soil variability, soil type, temperature and moisture. Experience and proper chemical placement have improved results of treatment.

Transmission of viruses through grapevine seed has been demonstrated in the case of false fanleaf virus (1820), peach rosette mosaic virus (1943) and Bulgarian latent virus (2090). Attempts to transmit fanleaf or fleck through grapevine seed gave negative results (1495, 1632, 1633, 1724). Several Nepoviruses are seedborne in weeds (1943, 1944).

Yellows diseases: mycoplasma, rickettsia-like organisms and bacterial diseases

Earlier this group of diseases were thought to be caused by viruses, because of their modes of transmission and responses. Now they may be separated into two groups — those that are caused by mycoplasma, for example Flavescence dorée and those caused by a bacterium, for example Pierce's disease. As research continues with these two groups of diseases, reports demonstrate vector-pathogen relationships and host-pathogen immune response, each of considerable interest in plant pathology. The application of advances in technology, isolation, culture, inoculation have contributed to the timely understanding of the nature of these two groups of diseases and opened new avenues to their control.

Diseases of these yellows types have been reported in the French Jura, Burgundy and Aube, in Switzerland, Italy and Germany. However, their etiology is not clear. — Bois noir is similar in many respects to Flavescence dorée but has a different etiology.

It has been suggested that a yellows disease present in vineyards of Western Germany (1887, 1888, 1889, 1988, 1989, 1990, 1991, 2018) and Greece (1989) with symptoms similar to those of Bois noir could be caused by rickettsia-like microorganisms, transmitted by *Xiphinema index*. An electron microscope study of the same yellows disease in Germany (1837) showed no microorganisms, but thread-like structures similar to virus particles in the phloem cell of infected grapevines.

In USSR, fleck and vein necrosis are considered to be caused by mycoplasma- or chlamidia-like microorganisms (1847, 1853, 1855, 1857).

In India, a little-leaf disease of grapevine is suspected to be caused by mycoplasmas (2019).

Diseases like Pierce's disease are known only in North America. However, a rickettsia-like organism is associated with infectious necrosis. It would be well to have a better understanding of this pathogen-host combination and its possible relationship to that of Pierce's disease.

References

Yellows diseases: mycoplasma-like organisms, rickettsia-like organisms, and bacteria

Yellows diseases

1503 — 1528 — 1580 — 1583 — 1588 — 1727 — 1728 — 1740 — 1760 — 1771
1837 — 1851 — 1884 — 1887 — 1888 — 1889 — 1892 — 1988 — 1989 — 1990
1991 — 2018 — 2082 — 2086.

Flavescence dorée

1445 — 1500 — 1501 — 1518 — 1528 — 1575 — 1576 — 1577 — 1579 — 1581
1582 — 1583 — 1584 — 1586 — 1587 — 1588 — 1589 — 1590 — 1722 — 1723
1772 — 1873 — 1874 — 1875 — 1892 — 2008 — 2052 — 2082 — 2086.

Bois noir

1525 — 1528 — 1585 — 1586 — 1588.

Pierce's disease

1466 — 1467 — 1468 — 1469 — 1610 — 1611 — 1697 — 1701 — 1739 — 1740
1741 — 1742 — 1743 — 1744 — 1745 — 1746 — 1747 — 1748 — 1785 — 1786
1799 — 1868 — 1869 — 1870 — 1871 — 1872 — 1912 — 1932 — 1933 — 1934
1936 — 1937 — 2071 — 2073.

Infectious necrosis

1649 — 1740 — 1803 — 1856 — 1862 — 2077 — 2128.

Other diseases

1839 — 1847 — 1853 — 1855 — 1857 — 2019.

Vectors and/or transmission

1518 — 1528 — 1575 — 1576 — 1579 — 1586 — 1588 — 1589 — 1590 — 1722
1868 — 1873 — 1875 — 1892 — 1932 — 1934 — 1935 — 2008 — 2009.

Vineyard amelioration and control of viroses

Most of the viroses and virus-like diseases of grapevines result in vine and vineyard degeneration. The factors that bear upon and influence the success of a program of amelioration and control of the viroses and virus-like diseases of the grapevine are: 1) the presence of the disease (the pathogen = inoculum), 2) the amount of disease (inoculum potential), and 3) the modes of spread, a) not spread naturally or rarely so or spread by man through nursery practices and distribution, b) air-borne or spread by insect vectors, and c) soil-borne or spread only in the soil by nematodes or fungi. Amelioration may be accomplished in either of two

ways and/or a combination program: 1) the "direct way"; selection within degenerating vineyards, testing performance and replanting with the best selected clones, and 2) control or the "clean health way"; selection in vineyards for high performance, rendering clones pathogen free, cleaning up soil and replanting with healthy selected clones (scion/rootstock combinations). Vineyards regenerated by the "direct way" will have relatively short productive lives, whereas those regenerated by the "clean health way" will have much longer useful lives. The two ways may be carried out at the same time in parallel programs. The direct way may be put into operation within some three to five years, while the clean health way may take up to ten or more years to function.

Vine selection on the basis of performance is the first step in either or both programs. It is well established that high vineyard performance (top quality and production) may be had by planting and/or replanting with high quality clones. In any selection program, it is good practice to select several clones and to replant with a random mixture of several of the high performance selected clones. Clones selected on a performance basis over three or more years are likely to be more stable. For example, some vines are known to produce good one year and poorly another. Selected clones of scion cultivars and rootstock should be planted in separate clonal plantings.

1) The "direct way" to vineyard amelioration

This may be termed the quick way; it requires the shortest time to become operative and costs are comparatively small.

Selections are made for trueness to variety, vigor, high quality, and production. Selected clones are propagated, increased in numbers and maintained in separate plantings. These clonal selected plantings serve as continuous test plantings for performance and as mother vines for replanting vineyards. Rootstocks, when used, should be as clean and free of viruses as possible. In areas and/or in some countries where soil-borne viruses are wide-spread and the vineyard soils contaminated with nematode vectors, selections are best made in several vineyards. Under these conditions, it is likely that all vines contain viruses. Some vines are usually found that perform much better than others. They have vigor and produce a good crop of fruit. Though the reasons why these few vines have only mild disease and good crops are not known, it appears that the more vigorous vine and its rootstock have a component of virus and/or viruses that cause only mild disease and "protect" the vine against the more severe forms of disease. When possible, it is simpler to force the rootstock of each clone (scion-rootstock combination) and to propagate the selected scion again on the same original rootstock. In case it is not possible to obtain the rootstock in this way, the clones should be propagated on clean (virus-free) rootstocks. Such rootstock lines are now available, they have been produced by thermotherapy, fully tested, and can be certified to be clean.

All selected clones and combination rootstocks should be grown in a separate multiplication block, (clonal mother block), in clean soil (free of the nematode vectors). Plants for replanting vineyards are propagated from the clonal mother vines. Rootstocks, each numbered to match each clone, are also grown in mother blocks.

Replanting infested vineyards: Vineyard soils can be sampled for the presence of nematode vectors. In most soils where vectors are found it is best to fallow the

soils (plant to no host crops) for some three or more years until vector nematode populations decline.

2) Control — the "clean health way" to vineyard amelioration

One of the cardinal principles in the control of a plant disease is to "plant clean propagating units in clean soil". The production of clean (pathogen-free) propagation material, planting in clean, vector-free soil, and the prevention of spread are thus of prime importance in control of the grapevine virus and virus-like diseases. Fortunately, most of the viruses that affect grapevines are soil-borne and the majority of virus-like diseases do not or rarely spread naturally. Control, therefore, has become directed to the soil-borne viroses. In this program, the selection of grapevine cultivar clones of high quality and high performance is one of the keys to success.

The program therefore requires: 1) the production of a number of clones of each of the desired grapevine cultivars and rootstocks, 2) maintaining clonal lines in good health, 3) performance testing clonal lines, 4) developing ways and means of cleansing infested soils, and 5) propagating and distribution of healthy materials. It will normally take some ten or more years to bring the program to maturity.

Healthy clones are produced after selection, by thermotherapy, and meristem culture, or combinations of meristem culture and thermotherapy, and may in the future be produced by way of protoplast (single cell) culture.

References

Selection, sanitation, preservation, propagation and certification

1389 — 1415 — 1418 — 1431 — 1433 — 1471 — 1475 — 1476 — 1477 — 1478
 1479 — 1481 — 1482 — 1483 — 1484 — 1485 — 1491 — 1492 — 1493 — 1494
 1496 — 1497 — 1498 — 1499 — 1502 — 1504 — 1523 — 1527 — 1530 — 1531
 1533 — 1536 — 1539 — 1540 — 1541 — 1542 — 1543 — 1544 — 1547 — 1563
 1569 — 1570 — 1572 — 1573 — 1594 — 1597 — 1608 — 1609 — 1612 — 1630
 1634 — 1636 — 1637 — 1642 — 1643 — 1647 — 1656 — 1657 — 1665 — 1668
 1678 — 1679 — 1681 — 1683 — 1692 — 1693 — 1700 — 1717 — 1731 — 1736
 1737 — 1749 — 1754 — 1755 — 1756 — 1787 — 1789 — 1797 — 1809 — 1823
 1831 — 1832 — 1834 — 1840 — 1876 — 1882 — 1891 — 1895 — 1903 — 1904
 1914 — 1916 — 1917 — 1919 — 1927 — 1940 — 1955 — 1961 — 1963 — 1971
 1972 — 2007 — 2013 — 2014 — 2017 — 2022 — 2023 — 2025 — 2029 — 2032
 2044 — 2047 — 2063 — 2069 — 2075 — 2076 — 2091 — 2092 — 2093 — 2095
 2097 — 2121 — 2122 — 2132 — 2137 — 2145 — 2146 — 2159.

Therapy (thermotherapy), meristem culture, shoot apex culture

1398 — 1455 — 1460 — 1472 — 1488 — 1489 — 1490 — 1526 — 1527 — 1528
 1537 — 1541 — 1660 — 1663 — 1670 — 1671 — 1672 — 1673 — 1692 — 1698
 1708 — 1769 — 1813 — 1842 — 1843 — 1844 — 1845 — 1877 — 1878 — 1879
 1880 — 1881 — 1893 — 1894 — 1908 — 1913 — 1960 — 2026 — 2033 — 2036
 2038 — 2039 — 2074 — 2076 — 2092 — 2096 — 2098 — 2110 — 2114 — 2119
 2120 — 2127 — 2141 — 2144 — 2148.

Performance

1450 — 1462 — 1527 — 1540 — 1548 — 1558 — 1628 — 1638 — 1751 — 1752
1788 — 1793 — 1795 — 2001 — 2006 — 2013 — 2070 — 2087 — 2106 — 2108
2117 — 2127.

Host-pathogen interaction: histology, cytology, physiology and biochemistry of disease, electron microscope studies

1390 — 1391 — 1424 — 1430 — 1438 — 1441 — 1442 — 1452 — 1486 — 1487
1667 — 1677 — 1718 — 1719 — 1803 — 1817 — 1837 — 1846 — 1848 — 1849
1850 — 1852 — 1854 — 1858 — 1865 — 1870 — 1885 — 1886 — 1910 — 1911
1992 — 1999 — 2000 — 2004 — 2005 — 2057 — 2074 — 2117 — 2133 — 2134.

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On évalue les proportions de Cicadelles devenues infectieuses en fonction des proportions observées de plantes devenues malades, lorsqu'il y a plusieurs cicadelles par plante. On évalue le nombre de particules infectieuses par millilitre en fonction des proportions de Cicadelles devenues malades.

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Des particules de type mycoplasme sont observées dans le liber des fèves inoculées par la Flavescence dorée et dans les cellules des vecteurs infectés. On ne les retrouve pas chez les fèves et les vecteurs sains témoins. Chez la vigne, les cellules libériennes apparaissent saines ou trop nécrosées. Les mycoplasmes n'ont pu être décelés que dans le cas de vignes inoculées artificiellement et qui ne montraient pas encore de symptômes.
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Les milieux liquides préconisés pour les mises en culture des mycoplasmes des animaux ne permettent pas la culture des agents pathogènes des jaunisses des plantes (Flavescence dorée en particulier). Il en est de même des milieux préconisés pour les «Mycoplasmes des végétaux». Ces milieux ont donné des résultats négatifs quelles que soient les multiples précautions prises. Nous préconisons l'utilisation exclusive de l'épreuve d'infectivité pour tester la survie ou la culture des agents pathogènes des jaunisses.
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