

Physics. — *Recovery and recrystallization viewed as processes of dissolution and movement of dislocations.* (Concluding part.) By W. G. BURGERS. (Laboratorium voor Physische Scheikunde der Technische Hoogeschool, Delft.) (Communicated by Prof. J. M. BURGERS.)

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III, 6. Occurrence of "stimulated" crystals.

We shall conclude this paper by discussing one other remarkable peculiarity deduced from the analysis of the forms of crystals in recrystallized aluminium plate. This refers to the presence of such "pointed" crystals as shown at *B* in fig. 18. From the analysis by SANDEE (62) and MAY (63) it follows that such crystals did not start to grow from their nucleus until another crystal (the "surrounding" crystal *A*), the growth of which

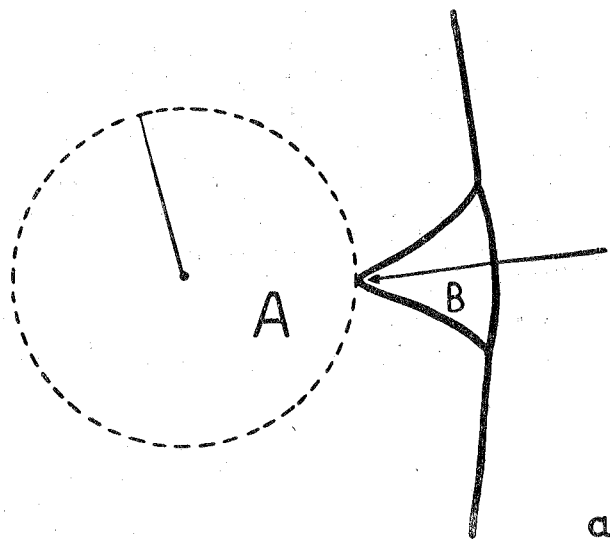


Fig. 19a. Formation of a "stimulated" crystal: Crystal *B* did not start to grow from its nuclear spot (indicated by the arrow) before crystal *A* had attained a size indicated by the circle with radius *r*.

had started previously, had come into contact with it in a way as described in fig. 19. In a discussion of this phenomenon (70) we have spoken in this connection of "stimulation" of growth of crystal *B* by crystal *A*. In order that the stimulated crystal obtains a measurable size, its rate of growth must be larger than that of the stimulating crystal. It was also found that both stimulating and stimulated crystals had a mutual orientation as found in spinel twins, that is they have a (111) plane and a (110) direction in common. As this plane and direction are those of preferential gliding in aluminium, it was suggested that the action of stimulation might be caused by something like a "discharge" of the stresses in the nucleus, which might take place by movement of dislocations along the common glide plane, at the moment that contact was established.

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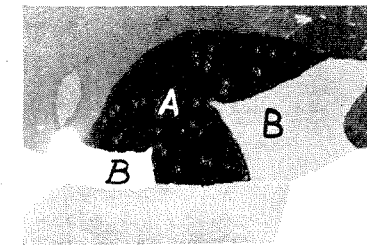
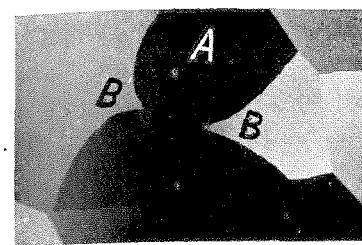


Fig. 18. Examples of stimulated crystals in recrystallized aluminium plate: the pointed crystals *B* have been "stimulated", in the sense as described in fig. 19a, by crystal *A*. Natural size.

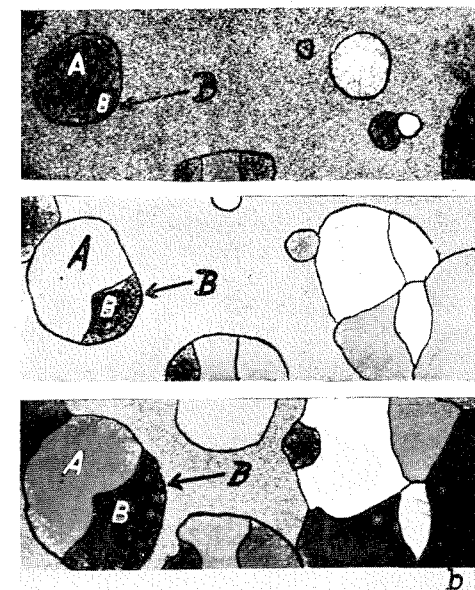


Fig. 19b. Formation of a "pointed" crystal in aluminium (cf. Fig. 19a). Natural size.

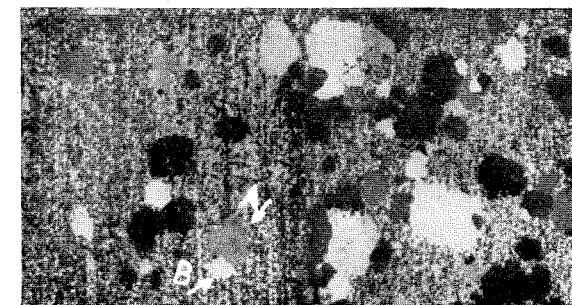


Fig. 21. Occurrence of "pointed" ("stimulated") crystals in the recrystallization experiments of ANDERSON and MEHL (10): it is highly probable that the pointed crystallite *B* (white on the reproduction) is "stimulated" by the growing crystal *A* (grey on the reproduction) (After ANDERSON and MEHL (10)). Magnif. about 7X.

The occurrence of stimulated crystals, which seems very common in recrystallized aluminium and which gives rise to most remarkable orientation relationships, is still a subject of closer investigation³³). Here, however, we mention this remarkable effect only in connection with some recent results, obtained by ANDERSON and MEHL (10) in their investigation of the recrystallization of thin aluminium sheet. From the increase of the

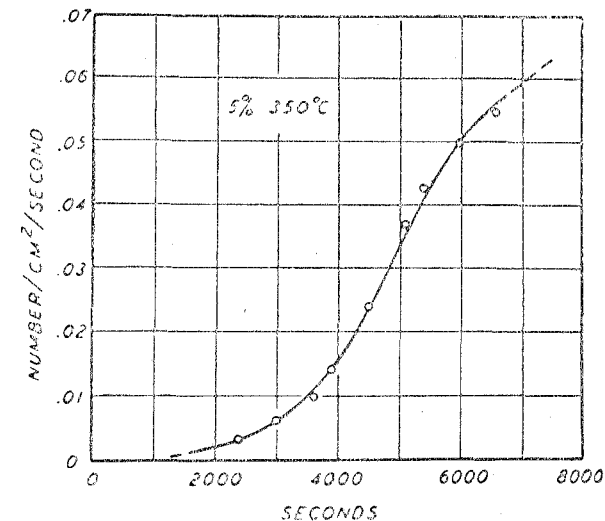


Fig. 20. Rate of nucleation N (number of grains formed per cm² of the unrecrystallized matrix per second) as a function of time of heating in isothermal recrystallization of thin aluminium sheet (350° C, 5.1% elongation). N is seen to increase in the course of the recrystallization process (after ANDERSON and MEHL (10)).

recrystallized fraction and of the number of new crystallites as a function of time of heating these authors deduced that, whereas the rate of growth of the new crystallites remained constant with time (in agreement with for example fig. 12), the rate of nucleation (that is the number of new grains formed per cm² of the unrecrystallized matrix per second) increased with time, at least in finegrained material³⁴). This is shown in fig. 20. With regard to this result ANDERSON and MEHL remark that the observed curve appears autocatalytic in nature and that such an auto-acceleration would appear to argue for an action at a distance; that is, that the occurrence and growth of one nucleus should transmit through the unrecrystallized matrix an accelerating stimulus to nucleation. Viewed in the light of the "stimulated" crystals discussed above, the actual occurrence of such an accelerating stimulus follows from the presence of "pointed" crystals of the type shown in fig. 18. That actually such crystals appear also in the recrystallized test-pieces of ANDERSON and MEHL is evident

³³) We are inclined to think that several other phenomena, observed while studying recrystallization, for example that of "secondary recrystallization" ("exaggerated graingrowth") and the formation of twins, is closely related to this stimulating action.

³⁴) In coarse-grained material a decrease was observed.

from fig. 21, which is taken from their paper: the small crystallite *B* is most probable an example of a stimulated crystal.

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LITERATURE.

1. W. G. BURGERS, Hdb. d. Metallphysik, Vol. III, 2 (Leipzig, Akad. Verlagsgesellschaft; Edwards Brothers, Ann. Arbor, Mich. 1944).
2. M. O. KORNFELD, Physik Z. Sowjetunion 7, 608 (1935).
3. U. DEHLINGER, Ann. Physik [5] 2, 749 (1929); Z. Metallkde 22, 221 (1930); 33, 16 (1941).
4. O. HAASE and E. SCHMID, Z. Physik 33, 413 (1925).
5. M. O. KORNFELD, Physik Z. Sowjetunion 6, 329 (1934).
6. F. KOREF, Z. Metallkde. 17, 213 (1925).
7. J. A. M. VAN LIEMPT, Rec. Trav. Chim. Pays-Bas 53, 941 (1934); Z. anorg. allg. Chem. 195, 366 (1931); see also W. G. BURGERS, Nature 159, 203 (1947).
8. A. KRUPKOWSKI and M. BALICKY, Rev. Métallurg. 36, 21 (1939); Ann. Acad. Sci. Techn. Varsovie 4, 270 (1937).
9. M. KORNFELD and W. PAWLOW, Physik Z. Sowjetunion 12, 301 (1937).
10. W. A. ANDERSON and R. F. MEHL, Am. Inst. Min. Met. Eng., Techn. Public. 1805 (1945).
11. H. G. MÜLLER, Z. Physik 96, 279 (1935).
- 11a. R. KARNOP and G. SACHS, Z. Physik 42, 283 (1927).
12. F. SAUERWALD, Z. Elektrochemie 37, 531 (1931).
- 12a. G. W. BRINDLEY, Proc. Leeds Philos. Soc. 4, 1, 167 (1941).
- 12b. G. MASING, private communication.
13. J. E. LENNARD JONES, Proc. Phys. Soc. 52, 38 (1940).
14. M. J. BUERGER, Z. Krist. 89, 195 (1934).
15. P. LACOMBE and L. BEAUJARD, Compt. Rend. 221, 414 (1945).
16. L. GRAF, Z. Elektrochem. 48, 181 (1942); Z. Physik 121, 73 (1943).
17. C. G. DARWIN, Phil. Mag. 27, 315, 675 (1914).
18. P. P. EWALD and M. RENNINGER, Pap. Disc. Intern. Conf. Physics (London 1934), Vol. II, p. 57; M. RENNINGER, Z. Krist. 89, 344 (1934).
- 18a. N. W. H. ADDINK, Nature 157, 764 (1946).
19. U. DEHLINGER and F. GISEN, Z. Physik. 35, 862 (1934); F. GISEN, Z. Metallkde. 27, 256 (1935).
20. A. SMEKAL, "Strukturempfindliche Eigenschaften der Kristalle", Hdb. d. Physik (Springer, Berlin) 2 Aufl. Vol. XXIV, 2 (1933).
21. W. L. BRAGG, Proc. Phys. Soc. 52, 105 (1940).
22. J. M. BURGERS, Proc. Phys. Soc. 52, 23 (1940); also Proc. Acad. Sci., Amsterdam, 42, 293 (1939).
23. W. L. BRAGG, Proc. Phys. Soc. 52, 54 (1940).
24. E. HERLINGER, Z. Krist. 92, 372, 380 (1935).
- 24a. M. BORN, J. Scient. Instr. 24 (1947) (Discussion X-Ray Conference).
25. F. ZWICKY, Helv. Phys. Acta 3, 269, 466 (1930); Proc. Nat. Acad. Sci. U.S.A. 15, 253, 816 (1929).
26. E. OROWAN, Pap. Disc. Intern. Conf. Physics (London), p. 81 (1934).
27. M. J. BUERGER, Z. Krist. 89, 242 (1934).
28. B. CHALMERS, Proc. Roy. Soc. London A 162, 120 (1937).
29. J. L. SNOEK, Z. Metallkde. 30, 94 (1938).
30. P. LACOMBE and L. BEAUJARD, Compt. Rend. 219, 66 (1944).
31. W. L. BRAGG, J. Scient. Instr. 19, 148 (1942); North-East Coast Inst. of Engineers and Shipbuilders 60, 299 (1944).
32. A. E. VAN ARKEL, Physica 5, 208 (1925).
33. W. P. DAVEY, Gen. Electric. Rev. 28, 588 (1925).
34. W. G. BURGERS and F. J. LEBBINK, Rec. Trav. Chim. Pays-Bas 64, 321 (1945); Proc. VIth International Congress of Applied Mechanics (Paris 1946); Nature 157, 47 (1946).
35. W. A. WOOD, Proc. Phys. Soc. 52, 110 (1940); Nature 143, 163 (1939).
36. U. DEHLINGER and A. KOCHENDÖRFER, Z. Metallkde. 31, 231 (1939); Z. Krist. 101, 134 (1939).
37. A. E. VAN ARKEL and W. G. BURGERS, Z. Physik, 48, 690 (1928); W. G. BURGERS, Proc. Phys. Soc. 52, 137 (1940).
38. H. D. MEGAW, H. LIPSON and A. R. STOKES, Nature 152, 20 (1943); 154, 145 (1944); J. Inst. Metals. 71, 279 (1945).
39. A. KOCHENDÖRFER, "Plastische Eigenschaften von Kristallen und metallischen Werkstoffen" (Berlin 1941) p. 119 et seq.
40. W. L. BRAGG, North-East Coast Inst. of Engineers and Shipbuilders 61, 25 (1945).
41. G. I. TAYLOR and H. QUINNEY, Proc. Roy. Soc. A 143, 307 (1934).
42. F. SEITZ and T. A. READ, J. Appl. Physics 12, 100 (1941).
43. J. S. KOEHLER, Phys. Rev. 60, 397 (1941).
44. C. BENEDICKS, Koll. Z. 91, 217 (1940).
45. E. OROWAN, Z. Physik 89, 634 (1934).
46. G. I. TAYLOR, Proc. Roy. Soc. A 145, 362 (1934).
47. W. LASCHKAREW and A. ALICHANIAN, Z. Krist. 80, 353 (1931).
48. N. A. BRILLIANTOW and J. W. OBREIMOW, Physik Z. Sowjetunion 3, 83 (1933).
49. W. G. BURGERS, Pap. and Disc. Intern. Conf. Physics (London 1934), Vol. II, p. 121.
50. U. DEHLINGER, Metallwirtsch. 12, 48 (1933).
51. C. S. BARNETT, "A New Microscopy and Its Potentialities", Am. Inst. Min. Met. Eng. Inst. Met. Div., Techn. Public. 1865 (1945).
52. M. COOK and C. MACQUARIE, Metals Techn. 5, (1938), Techn. Public. 974; M. COOK and T. L. RICHARDS, J. Inst. Metals 66, 1 (1940).
53. J. F. H. CUSTERS and G. W. RATHENAU, Physica 8, 759, 771 (1941).
54. M. KORNFELD and W. PAWLOW, Physik Z. Sowjetunion 6, 537 (1934).
55. J. A. COLLINS and C. H. MATHEWSON, Metals Techn. 7 (1940), Techn. Public. 1145.
56. M. KORNFELD and A. SCHAMARIN, Physik Z. Sowjetunion 11, 302 (1937).
57. E. OROWAN, Proc. Phys. Soc. 52, 8 (1940).
58. M. KORNFELD, Physik. Z. Sowjetunion 8, 533 (1935).
59. M. KORNFELD and F. RYBALKO, Physik. Z. Sowjetunion 12, 658 (1937).
60. R. KARNOP and G. SACHS, Z. Physik 60, 464 (1930).
61. M. KORNFELD and F. SAWIZKI, Physik Z. Sowjetunion 8, 528 (1935).
62. J. SANDEE, Physica 9, 741 (1942).
63. W. G. BURGERS and W. MAY, Rec. Trav. Chim. Pays-Bas 64, 5 (1945).
64. K. LONSDALE, Nature 151, 52 (1943); 153, 22, 433 (1944).
65. W. G. BURGERS and TAN KOEN HIOK, Physica 11, 353 (1946); Nature 157, 19 (1946); 160 (1947).
66. G. D. PRESTON, Proc. Roy. Soc. A 172, 116 (1939).
67. K. LONSDALE and H. SMITH, Proc. Roy. Soc. A 179, 8 (1941).
68. A. J. GUINIER, Proc. Phys. Soc. 57, 310 (1945).
69. R. KRONIG and J. J. ARLMAN; J. BOUMAN and L. L. VAN REYEN, Physica 10, 795 (1943); 12, 353 (1946).
70. W. G. BURGERS, Physica 9, 987 (1942); Nature 157, 76 (1946).

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