

***“This is a post-peer-review, pre-copyedit version of an article published as***

Larsson, A., Tollefsen, T.A., Løvvik, O.M. et al. (2019) A Review of Eutectic Au-Ge Solder Joints. *Metall and Mat Trans A* 50.

***The final authenticated version is available online at:***

<https://doi.org/10.1007/s11661-019-05356-0>

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

# 2 A Review of Eutectic Au-Ge Solder Joints



4  
5 ANDREAS LARSSON, TORLEIF A. TOLLEFSEN, OLE MARTIN LØVVIK,  
6 and KNUT E. AASMUNDTVEIT

7  
8 Gold-germanium (Au-Ge) joints have been part of the electronics industry since the birth of the  
9 solid state transistor. Today they find their role as a reliable joining technology, especially for  
10 high-temperature applications. This article is a literature study reviewing Au-Ge joints: Their  
11 uses, properties, material compatibility, application techniques, and performance characteris-  
12 tics. The review concludes that it is possible to create high-quality and very strong Au-Ge joints  
13 with a shear strength up to 150 MPa. They are stable and reliable, showing limited degradation  
14 after thousands of hours at high temperature and thousands of thermal cycles. Joints may be  
15 used in low-stress applications up to 300 °C.

16  
17 <https://doi.org/10.1007/s11661-019-05356-0>

18 © The Minerals, Metals & Materials Society and ASM International 2019

## 20 I. INTRODUCTION

21 RESEARCH on semiconductor materials and  
22 devices in the 1940s was primarily made on the  
23 semiconducting materials germanium (Ge) and silicon  
24 (Si). Gold was deposited onto germanium to solve a  
25 contact issue with an early prototype amplifier device  
26 (point contact transistor) by a research team led by  
27 William Bradford Shockley Jr. and Stanley Morgan at  
28 Bell Labs in 1947.<sup>[1]</sup> This device led the research team to  
29 the discovery of the *transistor effect*. William Bradford  
30 Shockley, John Bardeen, and Walter Houser Brattain  
31 were awarded the *Nobel prize* in physics in 1956 for their  
32 work on the transistor and the discovery of the  
33 transistor effect.<sup>[2]</sup> Thus, the Au-Ge contact was a  
34 central part of the birth of modern electronics. The first  
35 types of semiconducting transistors that became com-  
36 mercially available in the 1950s were also made from  
37 germanium. The more stable silicon replaced germa-  
38 nium in the 1960s, which changed the role of Au-Ge  
39 from providing a contact surface for interconnects (wire  
40 bonds) to a die-attach material joining dies to sub-  
41 strates. Eutectic Au-Ge bonding as a die-attach method  
42 was patented already in the early 1960s.<sup>[3,4]</sup> In the 1970s  
43 and 1980s, research on Au-Ge bonding focused more

44 towards ohmic contacts for gallium arsenide devices.<sup>[5-7]</sup>  
45 In the 1990s and especially post 2000, the high-temper-  
46 ature compatibility of Au-Ge joints has been thoroughly  
47 explored.<sup>[8-22]</sup> Environmental demands such as the  
48 RoHS directive have also lead to investigations on the  
49 Au-Ge system as a replacement for lead-based sol-  
50 ders.<sup>[23,24]</sup> The high material cost has most likely limited  
51 its applicability into volume mainstream, low-end elec-  
52 tronics devices.

53 The great majority of electronic devices and uses  
54 comprising Au-Ge joining technology has been explored  
55 with the utilization of a eutectic (or near eutectic)  
56 composition of the Au-Ge system. These alloys are  
57 today commonly used as a high-reliability, high-tem-  
58 perature compatible die-attach technology. The main  
59 motivation for this seems to be its high melting  
60 temperature at 361 °C<sup>[25]</sup> and the stable properties,  
61 corrosion, and thermal fatigue resistance, combined  
62 with excellent joint strength of the final joint. Typically,  
63 high-temperature applications with Au-Ge have aimed  
64 for use up to around 300 °C.<sup>[8,9,11,12,17,19,20,22,26-33]</sup> But,  
65 joints have also been explored for cryogenic tempera-  
66 tures down to around -170 °C.<sup>[15,26,32,35]</sup> Au-Ge joints  
67 have been evaluated as materials for die-attach purposes  
68 in numerous devices and configurations and various  
69 other uses in recent years. Applications include silicon  
70 carbide (SiC) power devices,<sup>[10,19,28,34-36]</sup> SiC  
71 diodes,<sup>[10,18-20,26,29,32,33,35]</sup> SiC dummy dies,<sup>[15,30,31]</sup> Si  
72 dummy dies,<sup>[8,9,11,12,17,22]</sup> microwave circuit,<sup>[27]</sup> MEMS  
73 device,<sup>[37]</sup> for wafer bonding,<sup>[37]</sup> creating nanowires,<sup>[38]</sup>  
74 forming ohmic contacts,<sup>[6,39]</sup> and hermetic seals<sup>[40,41]</sup> to  
75 mention some. Ceramic substrates (circuit boards) have  
76 most commonly been the substrate of choice in the  
77 evaluated systems. In particular, substrates of silicon  
78 nitride (Si<sub>3</sub>N<sub>4</sub>)<sup>[12,15,17,20,30,31,34-36]</sup> or alumina  
79 (Al<sub>2</sub>O<sub>3</sub>)<sup>[10-12,18,27,28]</sup> have been used. Other ceramic  
80 substrates include AlN<sup>[15,19]</sup> and BeO.<sup>[26,33]</sup> Both

ANDREAS LARSSON is with the Department of Applied Physics, TECHN I AS, 3184, Borre, Norway and also with the Departments of Materials and Micro-integration, University of South-Eastern Norway (USN), 3184, Borre, Norway. Contact e-mails: [anla7007@gmail.com](mailto:anla7007@gmail.com), [andreas.larsson@techni.no](mailto:andreas.larsson@techni.no), and [ala@usn.no](mailto:ala@usn.no) TORLEIF A. TOLLEFSEN is with the TEGma AS, 3015, Drammen, Norway. OLE MARTIN LØVVIK is with the Department of Sustainable Energy Technology, SINTEF Industry, 0373, Oslo, Norway. KNUT E. AASMUNDTVEIT is with the Department of Materials and Micro-integration, University of South-Eastern Norway (USN).

Manuscript submitted February 8, 2019.

	Journal : <b>MMTA</b>	Dispatch : <b>1-7-2019</b>	Pages : <b>10</b>
	PIPS No. : <b>5356</b>	<input type="checkbox"/> LE	<input type="checkbox"/> TYPESET
	MS Code :	<input type="checkbox"/> CP	<input type="checkbox"/> DISK

81 Cu<sup>[16,22]</sup> and Kovar<sup>[27]</sup> have also been used as substrates  
 82 (or lead frames).

## 83 II. MATERIAL PROPERTIES

84 Most materials properties of the Au-Ge system have  
 85 been evaluated for the eutectic (or near eutectic)  
 86 composition; 72 at. pct Au and 28 at. pct Ge, as shown  
 87 in the phase diagram of Figure 1. All data presented in  
 88 this review are based on this composition unless  
 89 otherwise specified. The properties of pure Au and Ge  
 90 have been thoroughly reported elsewhere and are not  
 91 part of the scope for this report. As shown from the  
 92 phase diagram of the Au-Ge system in Figure 1, only  
 93 three stable condensed phases exist; the solids  
 94 fcc\_A1 (Au) and diamond\_A4 (Ge), and a liquid (L).<sup>[42]</sup>  
 95 However, it should be mentioned that Tasci  
 96 *et al.* have reported the existence of a stable stoichiometric  
 97 phase, Au<sub>5</sub>Ge<sub>2</sub>, at lower temperatures<sup>[43]</sup> and  
 98 Maganin *et al.* identified a stable Au<sub>3</sub>Ge phase after  
 99 deposition of a Au film onto a Ge substrate.<sup>[44]</sup> These  
 100 phases will not be discussed further in this review. The  
 101 solubility of Ge in Au is up to 3.08 at. pct at the eutectic

temperature, 361 °C, and less than 1 at. pct at room  
 temperature.<sup>[25]</sup> The solubility of Au in Ge is negligi-  
 ble.<sup>[25]</sup> The chemical bonds between Au and Ge in  
 eutectic Au<sub>72</sub>Ge<sub>28</sub> have been identified to be covalent.  
<sup>[44-46]</sup> Eichhammer *et al.* illustrated the possibility to  
 significantly reduce the solidus intersecting the eutectic  
 point by using nanosized particles of 5 and 10 nm.<sup>[47]</sup>  
 Kryshstal *et al.* showed that for a bilayer of Au/Ge, the  
 Au film mass thickness needs to be larger than 0.2 nm  
 for a eutectic compound to form on the surface.<sup>[48]</sup> Near  
 eutectic compositions have an irregular lamellar type  
 microstructure, see Figure 2(a).<sup>[8,9,12,15,16,19,20,23,27,49]</sup>  
 Another common microstructure found in literature is  
 colonies of Ge dispersed in a Au matrix,<sup>[8,9]</sup> see  
 Figure 2(b). This second microstructure is especially  
 common after thermal aging which typically coarsens  
 the grain structure.<sup>[8,9,16,19,20]</sup> In contrast, Chidambaram  
*et al.* found that the microstructure of eutectic Au<sub>72</sub>Ge<sub>28</sub>  
 was refined after aging at 200 °C for three weeks.<sup>[23]</sup> No  
 further explanation to this rather contradictory result  
 was given. One explanation could be variations in  
 cooling rates during fabrication which could cause  
 different lamellar spacing.<sup>[50]</sup> Similarly, coarsened

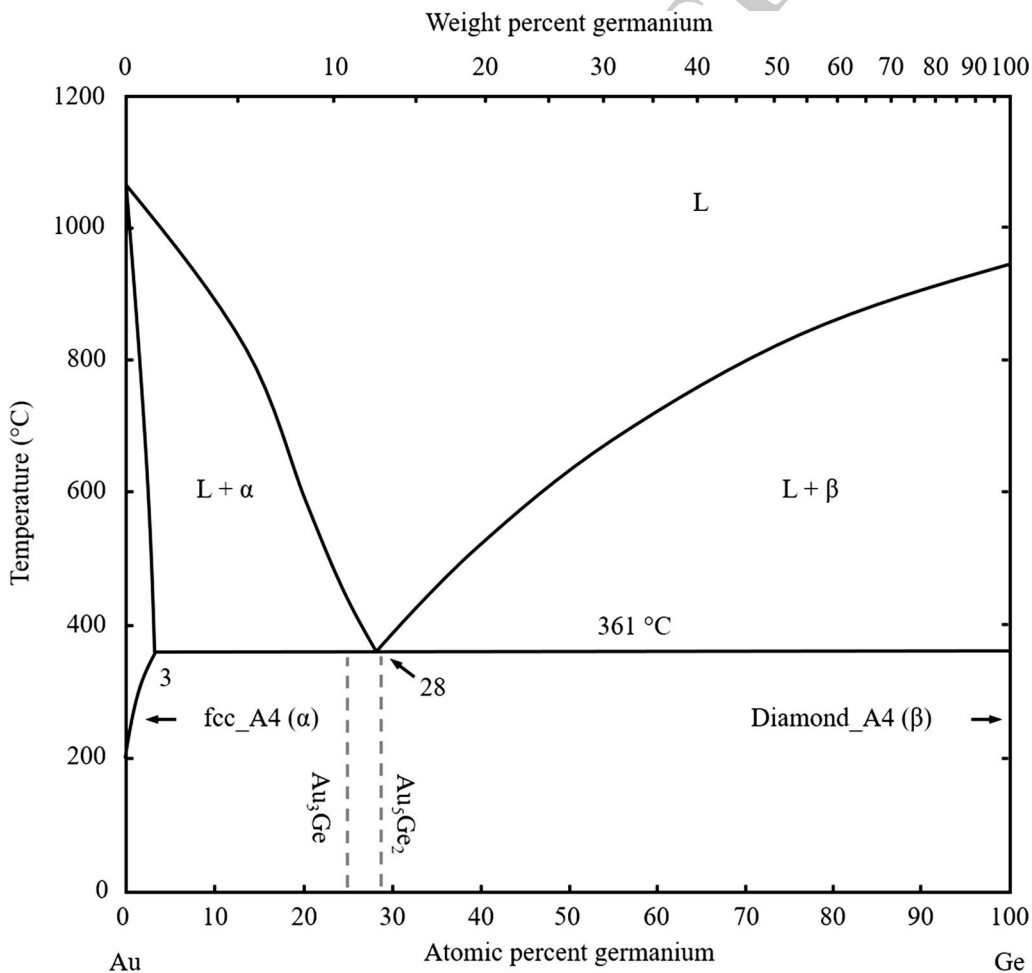


Fig. 1—The binary phase diagram of the Au-Ge system. The phase diagram was adapted from.<sup>[25,42-44]</sup> Note that the eutectic melting point varies slightly in the pertinent literature; 356 °C to 361 °C.<sup>[42,51]</sup>

125 microstructures can also be found in samples with  
 126 adjoining depletion layers, such as nickel (Ni).<sup>[9,16,19,20]</sup>  
 127 Typical static mechanical and thermal properties  
 128 found in literature are compiled in Table I. Values vary  
 129 somewhat between sources; thus, a range is provided  
 130 when appropriate. Discrepancies between the references  
 131 may originate from different setups during measure-  
 132 ment, purity of the samples, the morphology of the  
 133 different phases, heat treatment profiles, *etc.*

### 134 III. MATERIAL COMPATIBILITY

135 Material compatibility is crucial for any packaging  
 136 system. Joining dissimilar materials creates new phases,  
 137 *e.g.*, intermetallic compounds (IMC), and interfaces  
 138 between these different phases. They have dissimilar  
 139 properties, such as their coefficient of thermal expansion  
 140 and Young's modulus, which may cause high-stress  
 141 states in fabricated joints. Adjoined materials need to be  
 142 chemically stable and provide suitable mechanical, ther-  
 143 mal, and electrical performance for proper functionality  
 144 and reliability. The final microstructure and

145 composition may also depend on a wide variety of  
 146 parameters such as how the fabrication process was  
 147 carried out, *e.g.*, the deposition method used for the  
 148 metallization on dies and substrates, and temperature  
 149 profiles. Thus, care should be taken before concluding  
 150 on apparently similar systems. Further comparison  
 151 between different systems from different studies is even  
 152 more troublesome. This section presents general trends  
 153 and results extracted from the pertinent literature.  
 154 Juxtaposed materials systems evaluated in literature  
 155 are compiled below.

#### A. Nickel/Gold (ENIG)

156 Most commonly, eutectic Au-Ge have been used in  
 157 systems with a metallization scheme based on the  
 158 electroless nickel immersion gold (ENIG) sys-  
 159 tem.<sup>[8,9,11,12,15,19,20,22,26-28,31,33,35,36,49]</sup> In such systems,  
 160 Au-Ge typically reacts with Ni and forms Ge-Ni  
 161 intermetallic compounds (IMC). Lang *et al.* and  
 162 Egelkraut *et al.* reported the intermetallic phase to be  
 163 near the stoichiometric phases NiGe and Ni<sub>2</sub>Ge.<sup>[19,20]</sup>  
 164 Chidambaram *et al.* reported a Ni<sub>5</sub>Ge phase along with  
 165

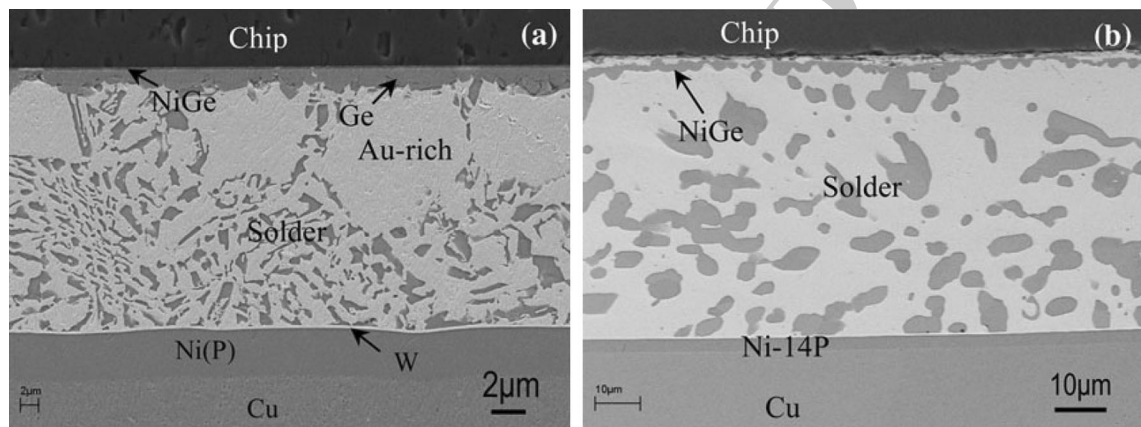


Fig. 2—Scanning electron microscope (SEM) image of cross-sections illustrating the microstructure of (a); a virgin joint and (b); a joint after heat treatment at 330 °C for 1000 h.<sup>[20]</sup> A coarsening of the microstructure is seen after thermal aging. Reprinted with permission.

Table I. Properties of Eutectic Au<sub>72</sub>Ge<sub>28</sub>

Property	Unit	Value	Source
Eutectic Composition	at. pct Ge	27.0 to 29.4	[25, 42]
Melting Point	°C	356 to 361 <sup>a</sup>	[25, 42, 47]
Young's Modulus	GPa	50 to 75 <sup>b</sup>	[18, 21, 52]
Poisson's Ratio		0.32	[21]
Yield Strength	MPa	100 to 240 <sup>b</sup>	[18, 52]
Shear Strength	MPa	220	[53]
Ultimate Strength	MPa	175 to 185 <sup>b</sup>	[52]
Elongation at Break	pct	10 to 50 <sup>b</sup>	[18]
Hardness	GPa	3.6	[8]
Coefficient of Thermal Expansion (CTE)	ppm/K	10.2 to 16.5 <sup>b</sup>	[52]
Thermal Conductivity	W/mK	44 to 44.4	[15, 52]
Electrical resistivity	μΩ cm	15 to 29	[32, 53]
Activation Energy—Creep	kJ/mol	11	[15]
Heat of Crystallization (a Ge)	J/mol	1.15	[54]

<sup>a</sup>Particle size dependent.

<sup>b</sup>Temperature dependent.

166 the NiGe phase. The growth rate of Ni<sub>2</sub>Ge was  
 167 measured to be up to a few nanometers per hour at  
 168 200 °C to 250 °C.<sup>[16]</sup> Godignon *et al.* evaluated the  
 169 electrical characteristics of a SiC diode joined with an  
 170 ENIG-like system and found it to be stable at temper-  
 171 atures between - 170 °C and 270 °C.<sup>[26,33]</sup> They also  
 172 found that the thermal resistance from junction to case  
 173 increased as a function of ambient temperature with  
 174 about 65 pct (0.6 K/W). The temperature dependence of  
 175 the joint strength was reported to show a nearly linear  
 176 decrease from about 115 MPa down to zero at the  
 177 melting point of eutectic Au-Ge.<sup>[15]</sup> Stress tests may be  
 178 found in References 8, 9, 19, 20, 26, 31, 33, 35, and 36.  
 179 The phase diagram of eutectic Au-Ge with Ni can be  
 180 found in Reference 27.

### 181 B. Copper

182 When Au-Ge is bonded directly to Cu, the compo-  
 183 nents react and create new intermetallic compounds  
 184 comprising Au-Cu-Ge.<sup>[8,9,16]</sup> Egelkraut *et al.* identified it  
 185 to be (Au,Cu)<sub>5</sub>Ge (the  $\xi$  phase) with varying Au-Cu  
 186 compositions.<sup>[16]</sup> They further measured the growth rate  
 187 to be about 5 to 10 nm per hour at 200 °C to 250 °C.  
 188 They also compared the results between Ni and Cu  
 189 metallization and found that Cu could create stronger  
 190 and more stable bonds than with the Ni metallization.  
 191 Between the IMC and the Cu layer, they further  
 192 observed a Au-Cu solid solution. The results indicate  
 193 an interdiffusion process, or material transport, of Au  
 194 and Cu through the joint, and that Ge does not interact  
 195 significantly with the Cu. The bond seems to change its  
 196 properties initially to finally stabilize at a new level when  
 197 exposed to high temperatures.<sup>[8,9]</sup> The IMC formation  
 198 likely causes the change. Related stress tests can be  
 199 found in References 8, 9, 18, and 21.

### 200 C. Silver

201 Lang *et al.* studied asymmetric systems with Ni/Ag  
 202 thin film on one side of the joint.<sup>[19,20]</sup> In one study, they  
 203 identified the fracture surface to the side of the joint  
 204 where Ni-Ge IMC were present. In another similar study,  
 205 they later showed that the fracture surface could be  
 206 moved to the device/joint interface by using tungsten  
 207 (W) as a diffusion barrier between Ge and Ni. Tanimoto  
 208 *et al.* showed that Ag in a thin film of Ni/Ag was  
 209 completely dissolved by the adjacent Au-Ge material.<sup>[31]</sup>  
 210 Both Egelkraut *et al.* and Drevin-Bazin *et al.* used  
 211 devices with Ag films, but no joint/Ag film-related  
 212 results were reported.<sup>[15,16]</sup> Related stress test may be  
 213 found in References 8, 9, 18, and 21.

### 214 D. Other Systems

215 A variety of materials systems combined with Au-Ge  
 216 have been evaluated in the pertinent literature, including  
 217 Al,<sup>[34]</sup> Ti/Ti-W,<sup>[17]</sup> W,<sup>[20,32]</sup> and glass.<sup>[17]</sup> Lang *et al.*  
 218 showed that it was possible to bond to an Al metaliza-  
 219 tion creating strong joints, > 50 MPa, and with  
 220 stable electrical performance results.<sup>[34]</sup> Long-term eval-  
 221 uation or other stress tests of that system was not

performed, and one might suspect that the Al-Au-Ge 222  
 system may form the well-known problematic Al-Au 223  
 IMCs such as Al<sub>2</sub>Au (purple plague) and Al<sub>2</sub>Au<sub>5</sub> (white 224  
 plague).<sup>[55]</sup> 225

## 226 IV. APPLICATION TECHNIQUES/PROCESS

227 Au-Ge joints have typically been created using var- 227  
 228 ious methods that are similar to common soldering 228  
 229 techniques. Eutectic preforms have been used fre- 229  
 230 quently.<sup>[11,12,15-18,27,28,31,35]</sup> Thin film techniques by 230  
 231 deposition and patterning have also been evaluated,<sup>[37]</sup> 231  
 232 along with solder balls<sup>[8,9]</sup> and pastes.<sup>[34]</sup> Bonding has 232  
 233 been performed in a vacuum, inert atmospheres, reduc- 233  
 234 ing atmosphere, flux, and air. 234

235 Recent process parameters that have been used in 235  
 236 literature are compiled in Table II. Avoiding atmo- 236  
 237 spheric oxygen seems crucial to be able to create strong 237  
 238 uniform joints. Cleanliness of bond surfaces also has a 238  
 239 strong impact on the bond quality. Regarding the 239  
 240 process peak temperature and time, it seems to be 240  
 241 enough to generate a liquid phase (eutectic) that has 241  
 242 enough time (seconds) to interdiffuse with the adjacent 242  
 243 bond surfaces to create strong bonds. Longer times or 243  
 244 higher temperatures affect the resulting microstructure 244  
 245 somewhat, but do not seem to have a direct impact on 245  
 246 the final bond quality. It has been demonstrated that it is 246  
 247 possible to create uniform joints with excellent coverage 247  
 248 and without significant voiding or microcracks inside 248  
 249 the joint.<sup>[11,12]</sup> 249

## 250 V. PERFORMANCE

### 251 A. Shear Strength

252 Shear strength tests are one of the most widely used 252  
 253 methods in literature for evaluating the bond quality. 253  
 254 Mechanical strength is often a more revealing measure 254  
 255 for bond quality than changes in electrical characteris- 255  
 256 tics of devices. Tanimoto and Matsui illustrated this 256  
 257 effect clearly in Reference 35. They measured a reduc- 257  
 258 tion in joint shear strength of several tens of percent 258  
 259 while the leakage current remained unchanged when 259  
 260 exposed to thermal cycling. A compilation of the shear 260  
 261 strength for virgin joints is presented in Figure 3. 261  
 262 Whenever possible, individual data points have been 262  
 263 extracted from the references. Extracted test results have 263  
 264 been assumed to have been carried out at 25 °C 264  
 265 whenever no specific information has been reported. 265  
 266 Shear test parameters are rarely disclosed in literature 266  
 267 and are thusly not compiled here. It is clear that the final 267  
 268 joint strength varies greatly, with a factor of more than 268  
 269 ten times, between reported devices (see results at 269  
 270 25 °C). The variation may originate from sample 270  
 271 preparation or simply from different system configura- 271  
 272 tions of the final assembly. Nonetheless, it is clear that it 272  
 273 is feasible to create very strong joints with eutectic 273  
 274 Au-Ge. It is also interesting to notice that the shear 274  
 275 strength at elevated temperatures, near the melting point 275  
 276 of the Au-Ge compound, is still significant. It should be 276

**Table II. Process Parameters Used to Fabricate Au-Ge Joints**

Peak Temperature (°C)	Preheat Temperature (°C)	Time at Peak Temperature (s)	Pressure	Atmosphere	Shear Strength (MPa)	Source
430	—	—	—	—	115	[15]
400	—	10	—	air	> 8	[28]
390	—	15	—	flux in air	22 to 34	[8,9]
—	—	—	—	N <sub>2</sub> AND VACUUM	—	[21]
> 363	—	300	2.1 MPa	vacuum or N <sub>2</sub>	—	[37]
430	—	120	—	vacuum	50 to 116	[34]
385	—	120 to 180	—	vacuum	26 to 52	[17]
410	—	—	—	vacuum	64 to 72	[20]
400	200	quick	—	red/vacuum/N <sub>2</sub>	128	[31]
—	—	—	> 0	red/vacuum	—	[26]
—	—	—	—	red/vacuum/N <sub>2</sub>	25	[16]
450	300/330	1800	> 0	red	78	[12]
—	—	—	—	red	240	[18]

Red: Reducing atmosphere (formic acid or H<sub>2</sub>).

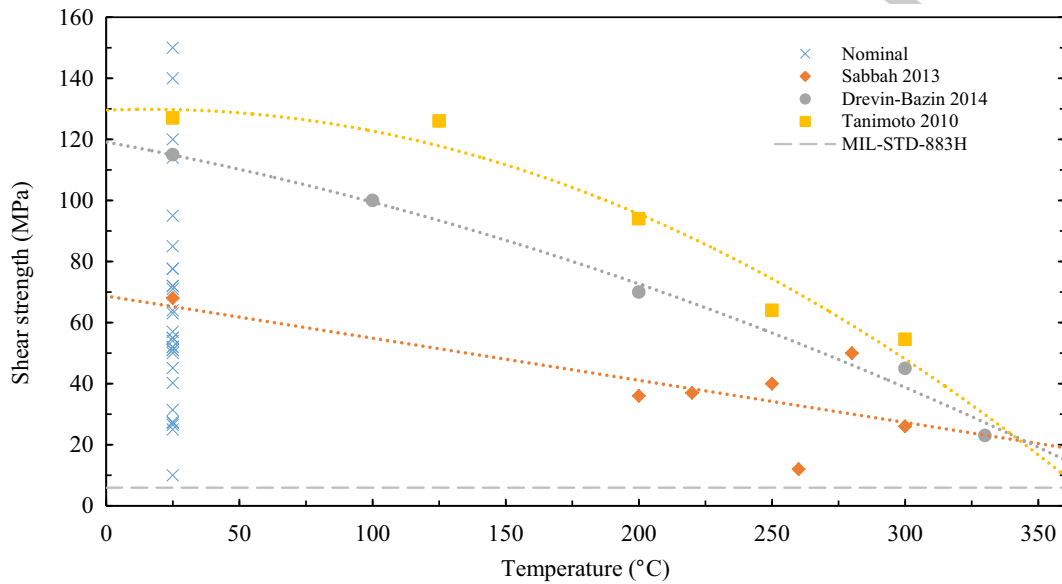


Fig. 3—Shear strength of virgin eutectic Au-Ge joints as a function of temperature for various devices.<sup>[8–12,15–17,19,20,22,28,31,34–36,49]</sup> Various samples tested at room temperature are compiled in the ‘Nominal’ series. Dashed lines are curve fits for each series. The strength requirement for solders, as stated in the US military standard (method 2019.8), is included here for Ref. [56].

277 pointed out that the reported strength results mirror the  
 278 weakest link in the tested system, not necessarily the  
 279 Au-Ge joint itself. Shear strength at cryogenic temper-  
 280 atures is yet to be evaluated.


281 **B. Electrical Evaluation**

282 Another way to characterize the bond quality is by  
 283 electrical evaluation. Different devices have been elec-  
 284 trically evaluated at temperatures from – 170 °C up to  
 285 330 °C.<sup>[10,19,29,32,35]</sup> The characteristics are typically  
 286 evaluated on a system level where the entire assembly  
 287 influences the results: the die, die-attach, wire bonds, *etc.*  
 288 Waveforms, current, voltage, and characteristic drift  
 289 have also been evaluated. The reported results typically  
 290 showed a limited degradation of the electrical perfor-  
 291 mance, and that this degradation could be directly

292 accredited to the joint itself. The thermal performance  
 293 has been evaluated at similar temperatures.<sup>[26]</sup> It was  
 294 found that the thermal resistance from junction to case  
 295 increased from 0.92 to 1.53 K/W.

296 **C. Stress Testing**

297 Stressing joints by exposing them to different loads is  
 298 a common way to evaluate their performance and  
 299 stability. Common techniques to stress devices include  
 300 thermal storage at elevated temperatures, thermal  
 301 cycling, and power loading. Different techniques stress  
 302 different parameters or mechanisms, *e.g.*, thermal stor-  
 303 age typically accelerates diffusion processes, thermal  
 304 cycling stimulates material fatigue, and power loads  
 305 excite migration mechanisms.

	Journal : <b>MMTA</b>	Dispatch : <b>1-7-2019</b>	Pages : <b>10</b>
	PIPS No. : <b>5356</b>	<input type="checkbox"/> LE	<input type="checkbox"/> TYPESET
	MS Code :	<input type="checkbox"/> CP	<input type="checkbox"/> DISK

306 In general, literature demonstrates a joint that is very  
 307 robust if given a proper design and applied within  
 308 certain load conditions. It can withstand very high  
 309 temperatures, near the melting point, as well as survive  
 310 wide thermal cycles. Electrical characterization results  
 311 also indicate a limited migration degradation when  
 312 stressed.

313 One interesting observation from this review comes  
 314 from an analysis of the combined results of Tanimoto  
 315 *et al.*,<sup>[35]</sup> Drevin-Bazin *et al.*,<sup>[15]</sup> and Msolli *et al.*<sup>[21]</sup>  
 316 Together, they point that the strength capacity of  
 317 eutectic Au-Ge joints significantly depends on the  
 318 mechanical load state. Drevin-Bazin *et al.* and Msolli  
 319 *et al.* showed this by studying the creep behavior.  
 320 Tanimoto *et al.* indirectly showed that fatigue failures  
 321 were reduced by improving their system for CTE  
 322 mismatch, thus reducing the stress state in the joint  
 323 during cycling. Note that this change might have shifted  
 324 the cycling characteristics, or failure mechanism,  
 325 between low cycle fatigue (LCF) and high cycle fatigue  
 326 (HCF). A detailed analysis of the joints would reveal  
 327 this.

#### 328 D. Thermal Storage

329 Figure 4 shows compiled thermal storage results.  
 330 There is a widespread between reported results. A  
 331 common denominator seems to be that proper diffusion  
 332 barriers between adjoined materials are key for strong  
 333 and stable joints. Results from Tanimoto *et al.* (250 °C  
 334 series at the top) show very strong joints with no  
 335 significant sign of degradation concerning high-temper-  
 336 ature exposure.<sup>[35]</sup> In an earlier report on a similar  
 337 system, they reported significant degradation of the joint  
 338 strength which may be accredited to poor initial joint  
 339 quality.<sup>[31]</sup> Other groups have shown similarly  
 340 stable trends for temperatures near the melting point

of eutectic Au-Ge.<sup>[11,17,20]</sup> Again, analyzing the results  
 from devices stored at 250 °C (top and bottom series in  
 Figure 4)<sup>[16,35]</sup> clearly illustrates how differently a joint  
 can behave in two different systems. The shear strength  
 differs by a factor of 10 between the two systems, even  
 though utilizing the same bonding technology. The  
 shear strength at 250 °C is reduced by about 30 pct after  
 500 hours at 300 °C.<sup>[9]</sup>

Among others, Godignon *et al.* have shown  
 stable electrical properties in their assembled SiC  
 diodes,<sup>[26,29,33]</sup> which indicate stable joint properties.  
 Degradation of electrical properties may typically not be  
 explicitly attributed to the joint. Note that joints  
 evaluated in non-optimized systems, *e.g.*, with improper  
 diffusion barriers,<sup>[29]</sup> show significant changes in electri-  
 cal properties. Electrical properties in various evaluated  
 systems may also be found in References 19 and 20.

#### E. Thermal Cycling

Au-Ge joints have been thermally cycled in a wide  
 variety of temperature ranges, from cryogenic temper-  
 atures down to -170 °C<sup>[26,29,32,33]</sup> and up to tempera-  
 tures as high as 325 °C.<sup>[17]</sup> Cycling rates have varied  
 between a few °C/min<sup>[10]</sup> and 40 °C/min<sup>[12,22,26,29,32,33]</sup>  
 with varying dwell times.

Tanimoto *et al.* showed initially that performance  
 degradation was significant due to thermal cycling by  
 observing the joint strength evolution.<sup>[31,36]</sup> The degra-  
 dation mechanism was coupled to an oxidation process  
 in cavities inside the bond. They later showed that by  
 optimizing the process with rigorous control of surface  
 cleanliness and optimizing the system CTE mismatch,  
 they could improve joint quality and both overall shear  
 strength and cycling performance significantly.<sup>[35]</sup> God-  
 ignon *et al.* indicated excellent thermal cycling perfor-  
 mance.<sup>[26,32,33]</sup> They inspected joints after exposure to

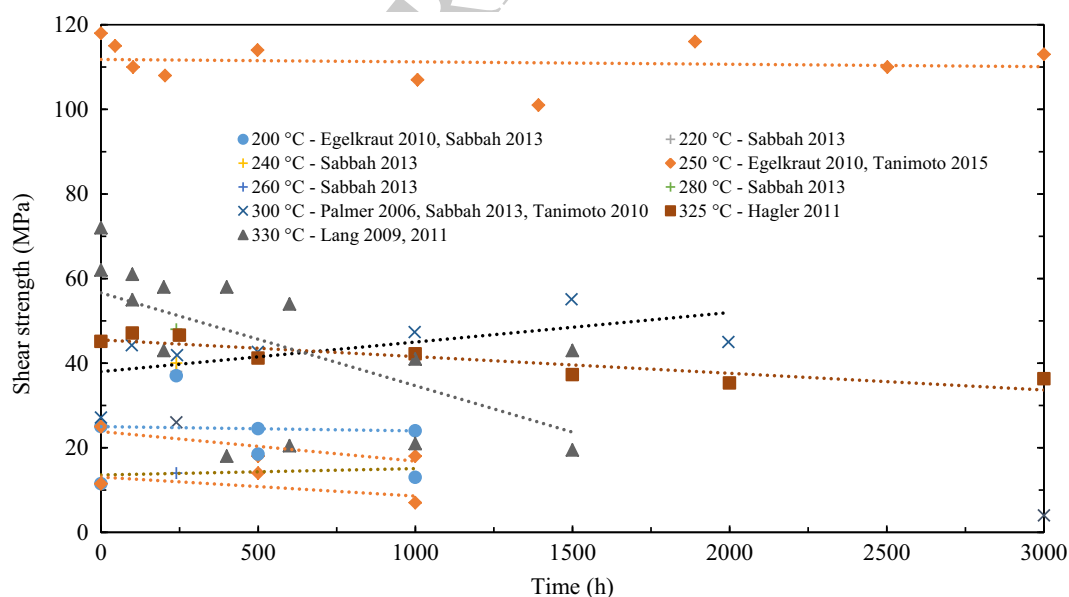


Fig. 4—Shear strength of eutectic Au-Ge joints in various devices after thermal storage between 200 °C and 330 °C.<sup>[8,9,11,12,16,17,19,20,28,31,35]</sup>  
 Dashed lines are curve fits for each series.

376 thousands of extended thermal cycles with a scanning  
 377 acoustic microscopy (SAM) technique, but no strength  
 378 evaluation was performed. In another study, it was  
 379 found that Au-Ge had limited thermal cycling capability  
 380 which was ascribed partly due to the joint stiffness.<sup>[22]</sup>  
 381 Zheng *et al.* tested a large joint (56.25 mm<sup>2</sup>) and found  
 382 that it has a shear strength of more than 17 MPa after  
 383 2000 cycles between 40 °C and 325 °C, at ± 10 °C/min  
 384 with a 5-min dwell time.<sup>[14]</sup> Compiling these results, one  
 385 may conclude that Au-Ge joints' thermal cycling capac-  
 386 ity can be very good. This assumes that the temperature  
 387 itself is not too close to the melting point,  $T_m$ , (at  
 388 homologous temperatures exceeding 0.9  $T_m$ ) and that  
 389 the mechanical stress state inside the joint is not very  
 390 high. A compilation of thermal cycling results found in  
 391 literature is presented in Figure 5.

392 Hutzler *et al.* used power cycles to thermally cycle  
 393 devices. They applied 15 to 18 A in short cycles that  
 394 created a temperature increase of the joint of 130 °C.  
 395 The maximum temperature in the joint was between  
 396 155 °C and 250 °C depending on the test scheme.<sup>[18]</sup>  
 397 Their system survived up to hundreds of thousands of  
 398 such cycles before the die-attach failed. Tanimoto *et al.*  
 399 power-cycled diodes between 35 °C and 200 °C and  
 400 found no strength degradation after tens of thousands  
 401 of cycles.<sup>[35]</sup> Thermally cycled devices (diodes), up to  
 402 4000 cycles between -170 °C and 270 °C, have been  
 403 characterized electrically without any significant  
 404 degradation.<sup>[26,29,32,33]</sup>

#### 405 F. Electrical Load

406 Various electrical loadings have been used to stress  
 407 devices assembled with Au-Ge joints. In a series of  
 408 reports, Godignon *et al.* have characterized SiC diodes  
 409 under various electrical loads combined with tempera-  
 410 ture loads.<sup>[26,29,32,33]</sup> The diodes were biased at 5 A  
 411 between 260 °C and 330 °C for up to 2000 hours and no

electrical degradation were found. They further investi- 412  
 gated their devices biased at -300 V and at 270 °C and 413  
 for 500 hours, again without any significant degrada- 414  
 tion. In the earlier reports, they observed drift behavior, 415  
 but that was accredited to the Schottky barrier. 416  
 Exchanging the barrier from Ni to W solved the 417  
 issue.<sup>[29]</sup> The results indicate good-quality joints. 418

#### G. Mechanical Load

419 Shear-loaded samples were prepared and tested by 420  
 Msolli *et al.* for strain rates in the order of magnitude 421  
 $\mu\text{m/s}$ , and mechanical loads of a few tens of MPa at 422  
 temperatures up to 300 °C.<sup>[21]</sup> They found that the joints 423  
 were fairly stable with limited dependence on displace- 424  
 ment and load rates. The joints showed a secondary 425  
 creep rate of about 3  $\mu\text{m/hour}$  when a 16.7 MPa load 426  
 was applied at 200 °C. At 300 °C and 16.7 MPa, 427  
 measurements showed tertiary behavior after a few 428  
 hours. Results also indicated a kinematic hardening of 429  
 the material. Similar experiments were performed by 430  
 Drevin-Bazin *et al.*<sup>[15]</sup> Unlike Msolli *et al.* they did not 431  
 find a secondary stage. The joint showed a clear tertiary 432  
 stage for high stress. In general, they found that the 433  
 creep is thermally activated and stress-dependent. 434

## 435 VI. COMPARISON WITH OTHER 436 TECHNOLOGIES

437 Egelkraut *et al.* showed that Au-Ge joints had better 438  
 aging characteristics than lead-rich Pb-Sn joints.<sup>[16]</sup> 439  
 Navarro *et al.* showed that Pb-Sn-Ag joints have a 440  
 better thermal cycling capacity than Au-Ge joints.<sup>[22]</sup> 441  
 Nevertheless, the quality of the produced Au-Ge joints 442  
 varied greatly. Studies have compared Au-Ge joints with 443  
 other binary Au-based alloys. It has been shown that 444  
 Au-Sn alloys are more corrosion-resistant than Au-Ge

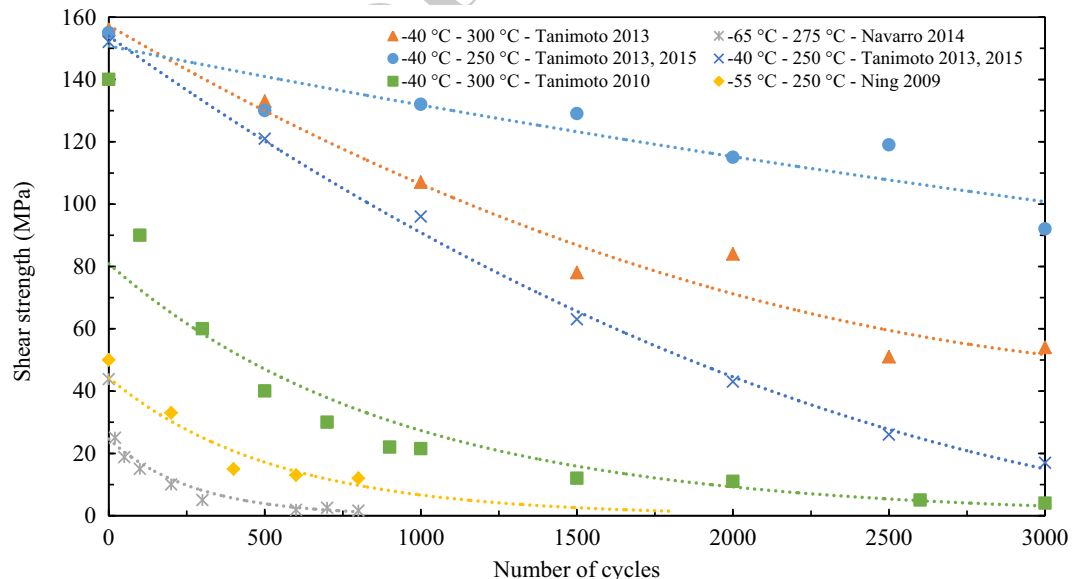


Fig. 5—Shear strength as a function of thermal cycles for various cycling regimes.<sup>[10,22,30,31,35,36]</sup> Dashed lines are curve fits for each series.



**Table III. Comparison Between Three High-Temperature Compatible Die-Attach Technologies; Eutectic Au-Ge, TLP/SLID, and Ag Sintering**

Process	Advantages	Disadvantages
Eutectic Au-Ge Soldering	simple and fast soldering type process scheme low bond line pressure easily repaired by desoldering tolerates rough and irregular bond surfaces self-aligning fluxless thermally stable at very high homologous temperatures ( $0.9 T_H$ ) excellent joint quality without significant voiding very strong joints (up to 150 MPa)	high process temperature operation temperature limited by eutectic melting point, $T_O < T_P$ oxygen-free atmosphere required during fabrication expensive materials no paste commercially available
TLP/SLID	high thermal joint stability ( $T_O > T_P$ ) relatively low bond line pressure (0.2 to 5 MPa) tolerates some surface roughness tolerable to faying surface oxide fluxless similar material properties as the base material in bond surfaces good joint quality very strong joints (up to 150 to 200 MPa)	requires flat and well-aligned bond surfaces time consuming (hours to days) lack of reparability formation of thick IMCs which reduces strength and ductility expensive materials
Ag Sintering	monometallic joints thermally stable joints (typically, $T_O = \text{low } T_H$ ) excellent electrical and thermal material properties	high process temperature high bond line pressure (up to tens of MPa) somewhat time consuming (approx. 30 to 60 min) poor joint quality—porosity (approx. 20 pct) Ag migration relatively low joint strength (approx. 20 to 30 MPa)

$T_P$ : process temperature,  $T_O$ : operation temperature,  $T_H$ : Homologous temperature, *IMC*: intermetallic compound.

Author Proof

445 alloys.<sup>[57]</sup> Lang *et al.* bonded Au-Ge and Au-Sn to Au  
 446 stud bumps on Al pads and found mechanical and  
 447 electrical characteristics of the produced joints to be  
 448 similar.<sup>[34]</sup> Chidambaram *et al.* compared Au-Ge and  
 449 Au-Si joints and found that the mechanical properties  
 450 were similar.<sup>[8,9]</sup> Aging tests showed that the shear  
 451 strength of the Au-Ge joints were about twice as strong  
 452 as similar Au-Si joints at room temperature and at  
 453 250 °C. Au-Ge joints have also been compared with  
 454 Au-In joints.<sup>[15]</sup> The Au-In joints showed better creep  
 455 behavior and higher strength at elevated temperatures as  
 456 they did not degrade as fast as Au-Ge. Nevertheless, at  
 457 300 °C, the two joints showed similar strength capacity  
 458 since the initial strength of the Au-Ge joints was higher.  
 459 Au-In (with roughly 30 to 67 at. pct In) and eutectic  
 460 Au-Ge melt at between 450 °C and 500 °C and 360 °C,  
 461 respectively. Thus, at 300 °C, the homologous temper-  
 462 ature for the two tested joints are 0.75 and 0.90,  
 463 respectively. The homologous temperature is the applied  
 464 temperature divided by the melting temperature, mea-  
 465 sured in Kelvin. It should also be noted here that the  
 466 Au-In preforms and Au-Ge preforms used in that study  
 467 had different thicknesses.

#### 468 A. High-Temperature Die-Attach Technologies

469 Since eutectic Au-Ge joints are commonly used in  
 470 high-temperature high-reliability applications, it is inter-  
 471 esting to compare it with other high-temperature com-  
 472 patible die-attach technologies such as transient liquid  
 473 phase (TLP) or solid-liquid interdiffusion (SLID)

bonding,<sup>[15,58–68]</sup> and silver sintering.<sup>[12,16,18,22,28,69,70]</sup>  
 Table III shows a comparison between eutectic Au-Ge,  
 TLP/SLID, and Ag sintering.

## 477 VII. RECOMMENDATIONS

478 Based on this literature study and the authors'  
 479 experience, the following recommendations can be made  
 480 for eutectic Au<sub>72</sub>Ge<sub>28</sub> joints:

- 481 • It is important to have a suitable diffusion barrier  
 482 between the joint and the adjoining layers to avoid  
 483 changes in the composition and microstructure of  
 484 the joint. Reaction with an adjoining Ni layer causes  
 485 Ge-Ni IMCs to form. These IMCs are stable at  
 486 high-temperature but seem to restrict the thermal  
 487 cycling performance. Tungsten (W) is an effective  
 488 barrier between Ni and the joint.
- 489 • The CTE mismatch between joined components  
 490 should be minimized to avoid fatigue issues during  
 491 thermal cycling. Alternatively, a mechanical absorp-  
 492 tion layer, *e.g.*, Au, could be used to reduce the stress  
 493 state inside the joint.
- 494 • Fabrication should be performed in an oxygen-free  
 495 atmosphere.
- 496 • Joints should not be used at temperatures above  
 497 300 °C as the shear strength of eutectic joints drops  
 498 rapidly at very high homologous temperatures  
 499 ( $T_H \approx 0.9$ ).

501 A compilation of literature shows that it is possible to  
 502 form high-quality joints using eutectic Au<sub>72</sub>Ge<sub>28</sub>. The  
 503 joints may have excellent thermomechanical properties  
 504 and are very stable at temperatures as high as 300 °C.  
 505 The shear strength may be up to 150 MPa at room  
 506 temperature and around 50 MPa at 300 °C. This  
 507 enables them for use in a wide variety of high-temper-  
 508 ature applications. To create such high-quality joints, it  
 509 is crucial to design the systems so that the stress state  
 510 inside the joint is limited and that fabrication is done in  
 511 an oxygen-free atmosphere. The main disadvantages are  
 512 material cost and unavailability of a commercial eutectic  
 513 paste. Thus, eutectic Au<sub>72</sub>Ge<sub>28</sub> is a suitable technology  
 514 for high-temperature, high-reliability, and high-end  
 515 applications.

516

517

## ACKNOWLEDGMENTS

518 We want to acknowledge the Norwegian Research  
 519 Council for supporting this project (Project No.:  
 520 244915).

521


## REFERENCES

- 522 1. AT&T Archives and History Center: *AT&T Archives Genesis of the*  
 523 *Transistor*, AT&T Tech Channel, New York, 1965.
- 524 2. The Nobel Prize in Physics 1956: [http://www.nobelprize.org/nobel\\_](http://www.nobelprize.org/nobel_prizes/physics/laureates/1956/)  
 525 [prizes/physics/laureates/1956/](http://www.nobelprize.org/nobel_prizes/physics/laureates/1956/). Accessed 13 November 2018.
- 526 3. United States Patent Office, 3,025,439, 1962.
- 527 4. United States Patent Office, 3,200,490, 1965.
- 528 5. K. Nishitani, O. Ishihara, H. Sawano, T. Ishii, S. Mitsui, and H.  
 529 Miki: *Jpn. J. Appl. Phys.*, 1976, vol. 16, pp. 93–97.
- 530 6. M.S. Islam and P.J. McNally: *Microelectron. Eng.*, 1998, vol. 40,  
 531 pp. 35–42.
- 532 7. A.A. Iliadis, J.K. Zahurak, T. Neal, and W.T. Masselink: *J.*  
 533 *Electron. Mater.*, 1999, vol. 28, pp. 944–48.
- 534 8. V. Chidambaram, H.B. Yeung, and G. Shan: *Proc. Int. Symp.*  
 535 *Phys. Fail. Anal. Integr. Circuits, IPFA*, [https://doi.org/10.1109/](https://doi.org/10.1109/ipfa.2012.6306308)  
 536 [ipfa.2012.6306308](https://doi.org/10.1109/ipfa.2012.6306308).
- 537 9. V. Chidambaram, H.B. Yeung, and G. Shan: *J. Electron. Mater.*,  
 538 2012, vol. 41, pp. 2107–17.
- 539 10. P. Ning, R. Lai, D. Huff, F. Wang, K.D.T. Ngo, V.D. Immanuel,  
 540 and K.J. Karimi: *IEEE Trans. Power Electron.*, 2010, vol. 25,  
 541 pp. 16–23.
- 542 11. M.J. Palmer, R.W. Johnson, and B.H. Ecedept: *Int. High Temp.*  
 543 *Electron.*, Santa Fe, NM, 2006.
- 544 12. W. Sabbah, S. Azzopardi, C. Buttay, R. Meuret, and E. Woigard:  
 545 *Microelectron. Reliab.*, 2013, vol. 53, pp. 1617–21.
- 546 13. V.R. Manikam and K.Y. Cheong: *Compon. Packag. Manuf.*  
 547 *Technol. IEEE Trans.*, 2011, vol. 1, pp. 457–78.
- 548 14. P. Zheng: Ph.D. dissertation, Dept. Electr. Comput. Eng., Auburn  
 549 Univ., Auburn, AL, 2010.
- 550 15. A. Drevin-Bazin, F. Lacroix, and J.F. Barbot: *J. Electron. Mater.*,  
 551 2014, vol. 43, pp. 695–701.
- 552 16. S. Egelkraut, L. Frey, M. Knoerr, and A. Schletz: *IEEE 12th Proc.*  
 553 *Electron. Packag. Technol. Conf. (EPTC)*, Singapore, 2010, pp.  
 554 660–67.
- 555 17. P. Hagler, P. Henson, and R.W. Johnson: 2011.
- 556 18. A. Hutzler, A. Tokarski, S. Kraft, S. Zischler, and A. Schletz:  
 557 *IEEE Electron. Compon. Technol. Conf.*, Orlando, FL, 2014, pp.  
 558 1700–06.
- 559 19. F.L.F. Lang, S. Tanimoto, H. Ohashi, and H. Yamaguchi: *2009*  
 560 *Eur. Microelectron. Packag. Conf.*, 2009, pp. 3–7.
20. F. Lang, H. Yamaguchi, H. Ohashi, and H. Sato: *J. Electron.*  
 561 *Mater.*, 2011, vol. 40, pp. 1563–71.
21. S. Msolli, O. Dalverny, J. Alexis, and M. Karama: *Integr. Power*  
 562 *Electron. Syst. (CIPS)*, 2010 6th Int. Conf., 2010, pp. 16–18.
- 563 22. L.A. Navarro, X. Perpina, P. Godignon, J. Montserrat, V. Banu,  
 564 M. Vellvehi, and X. Jorda: *IEEE Trans. Power Electron.*, 2014,  
 565 vol. 29, pp. 2261–71.
- 566 23. V. Chidambaram, J. Hald, and J. Hattel: *J. Alloys Compd.*, 2010,  
 567 vol. 490, pp. 170–79.
- 568 24. V. Chidambaram, J. Hald, and J. Hattel: *Microelectron. Reliab.*,  
 569 2009, vol. 49, pp. 323–30.
- 570 25. H. Okamoto and T.B. Massalski: *Bull. Alloy Phase Diagr.*, 1984,  
 571 vol. 5, pp. 601–10.
- 572 26. P. Godignon, X. Jorda, M. Vellvehi, X. Perpina, V. Banu, D.  
 573 Lopez, J. Barbero, P. Brosselard, and S. Massetti: *IEEE Trans.*  
 574 *Ind. Electron.*, 2011, vol. 58, pp. 2582–89.
- 575 27. L. Ma, X. Huang, and J. Zha: *Int. Conf. Electron. Packag. Technol.*,  
 576 2013, pp. 946–49.
- 577 28. R. Kisiel and Z. Szczepański: *Microelectron. Reliab.*, 2009, vol. 49,  
 578 pp. 627–29.
- 579 29. E. Maset, E. Sanchis-Kilders, J.B. Ejea, A. Ferreres, J. Jordán, V.  
 580 Esteve, P. Brosselard, X. Jordà, M. Vellvehi, and P. Godignon: *IEEE*  
 581 *Trans. Device Mater. Reliab.*, 2009, vol. 9, pp. 557–62.
- 582 30. S. Tanimoto, K. Watanabe, H. Tanisawa, K. Matsui, and S. Sato:  
 583 *Electrochem. Soc. Meet., 224th*, The Electrochemical Society, San  
 584 Francisco, CA, 2013, p. 1.
- 585 31. S. Tanimoto, K. Matsui, Y. Murakami, H. Yamaguchi, and H.  
 586 Okumura: in *IMAPS Int. Conf. High Temp. Election. (HiTEC)*,  
 587 IMAPS, Albuquerque, NM, 2010, pp. 32–9.
- 588 32. V. Banu, P. Godignon, X. Jorda, M. Vellvehi, J. Millan, P.  
 589 Brosselard, D. Lopez, and J. Barbero: *Proc. Int. Semicond. Conf.*  
 590 *CAS*, 2010, vol. 2, pp. 397–400.
- 591 33. P. Godignon, X. Jorda, V. Banu, M. Vellvehi, J. Millan, P.  
 592 Brosselard, D. Lopez, and J. Barbero: *Power Semicond. Devices*  
 593 *& IC's (ISPSD)*, 2010 22nd Int. Symp., 2010, pp.  
 594 351–54.
- 595 34. F. Lang, H. Nakagawa, and H. Yamaguchi: *Gold Bull.*, 2013,  
 596 vol. 47, pp. 109–18.
- 597 35. S. Tanimoto and K. Matsui: *IEEE Trans. Electron Devices*, 2015,  
 598 vol. 62, pp. 258–69.
- 599 36. S. Tanimoto, H. Tanisawa, K. Watanabe, K. Matsui, and S. Sato:  
 600 *Mater. Sci. Forum*, 2013, vols. 740–742, pp. 1040–43.
- 601 37. R. Chanchani, C.D. Nordquist, R.H. Olsson, T. Peterson, R. Shul,  
 602 C. Ahlers, T.A. Plut, and G.A. Patrizi: *Proc. Electron. Components*  
 603 *Technol. Conf.*, 2011, pp. 1604–09.
- 604 38. E.J. Schwalbach and P.W. Voorhees: *Nano Lett.*, 2008, vol. 8,  
 605 pp. 3739–45.
- 606 39. Y.-C. Shih, M. Murakami, E.L. Wilkie, and A.C. Callegari: *J.*  
 607 *Appl. Phys.*, 1987, vol. 62, pp. 582–90.
- 608 40. M. Pecht: *Integrated Circuit, Hybrid, and Multichip Module*  
 609 *Package Design Guidelines: A Focus on Reliability*, Wiley, New  
 610 York, 1994.
- 611 41. M. Schwartz: *Soldering - Understanding the Basics*, ASM Inter-  
 612 national, Materials Park, 2014.
- 613 42. J. Wang, C. Leinenbach, and M. Roth: *J. Alloy. Compd.*, 2009,  
 614 vol. 481, pp. 830–36.
- 615 43. E.S. Tasci, M.H.F. Sluiter, A. Pasturel, and N. Jakse: *Phys. Rev.*  
 616 *B*, 2010, vol. 81, pp. 1–3.
- 617 44. L. Magagnin, R. Maboudian, and C. Carraro: *J. Phys. Chem. B*,  
 618 2002, vol. 106, pp. 401–07.
- 619 45. D.G. Popescu and M.A. Husanu: *Rapid Res. Lett.*, 2013, vol. 7,  
 620 pp. 274–77.
- 621 46. D.G. Popescu and M.A. Husanu: *Thin Solid Films*, 2014, vol. 552,  
 622 pp. 241–49.
- 623 47. Y. Eichhammer, J. Roeck, N. Moelans, F. Iacopi, B. Blanpain,  
 624 and M. Heyns: *Arch. Met. Mater.*, 2008, vol. 53, pp. 1133–39.
- 625 48. A.P. Kryshal, R.V. Sukhov, and A.A. Minenkov: *J. Alloys*  
 626 *Compd.*, 2012, vol. 512, pp. 311–15.
- 627 49. V. Chidambaram, E.P.J. Rong, G.C. Lip, and M.W.D. Rhee:  
 628 *Electron. Packag. Technol. Conf.*, IEEE, Singapore, 2013, pp.  
 629 202–07.
- 630 50. F.C. Campbell: *Phase Diagrams - Understanding the Basics*, ASM  
 631 International, Materials Park, 2012.
- 632 51. R.P. Elliott and F.A. Shunk: *Bull. Alloy Phase Diagr.*, 1980, vol. 1,  
 633 pp. 51–54.
- 634 635

<p>636 637 638 639 640 641 642 643 644 645 646 647 648 649 650 651 652 653 654 655 656 657</p>	<p>52. D. Olsen and H. Berg: <i>IEEE Trans. Compon. Hybrids Manuf. Technol.</i>, 1979, vol. 2, pp. 257–63.</p> <p>53. M.F. Sousa, S. Riches, C. Johnston, and P.S. Grant: <i>High Temp.</i>, 2010, pp. 1–6.</p> <p>54. Z.W. Chen, J.K.L. Lai, and C.H. Shek: <i>J. Phys. D</i>, 2006, vol. 39, pp. 4544–48.</p> <p>55. <i>Materials for Advanced Packaging</i>, 1st ed., D. Lu, and C.P. Wong, eds., <i>Materials for Advanced Packaging</i>, Springer, New York, 2009.</p> <p>56. <i>MIL-STD-202G</i>, 2002.</p> <p>57. V. Chidambaram, J. Hald, R. Ambat, and J. Hattel: <i>JOM</i>, 2009, vol. 61, pp. 59–65.</p> <p>58. G.O. Cook and C.D. Sorensen: <i>J. Mater. Sci.</i>, 2011, vol. 46, pp. 5305–23.</p> <p>59. W.F. Gale and D.A. Butts: <i>Sci. Technol. Weld. Join.</i>, 2004, vol. 9, pp. 283–300.</p> <p>60. W.D. MacDonald and T.W. Eagar: <i>Annu. Rev. Mater. Sci.</i>, 1992, vol. 22, pp. 23–46.</p> <p>61. W.D. MacDonald and T.W. Eagar: <i>Met. Sci. Join.</i>, 1992, pp. 93–100.</p> <p>62. W.D. MacDonald and T.W. Eagar: <i>Metall. Mater. Trans. A</i>, 1998, vol. 29A, pp. 315–25.</p>	<p>63. L. Bernstein: <i>J. Electrochem. Soc.</i>, 1966, vol. 113, pp. 1282–88.</p> <p>64. L. Bernstein and H. Bartholomew: <i>Trans. Metall. Soc. AIME</i>, 1966, vol. 236, pp. 405–12.</p> <p>65. T.A. Tollefsen, A. Larsson, O.M. Løvvik, and K. Aasmundtveit: <i>Metall. Mater. Trans. B</i>, 2012, vol. 43B, pp. 397–405.</p> <p>66. T.A. Tollefsen, O.M. Løvvik, K. Aasmundtveit, and A. Larsson: <i>Metall. Mater. Trans. A</i>, 2013, vol. 44A, pp. 2914–16.</p> <p>67. K.E. Aasmundtveit, T.-T. Luu, H.-V. Nguyen, A. Larsson, and T.A. Tollefsen: in <i>Intermetallic Compounds - Formation and Applications</i>, M. Aliofkhazrai, ed., IntechOpen, London, 2018, 1st ed., pp. 43–72.</p> <p>68. T.T. Luu, N. Hoivik, K. Wang, K.E. Aasmundtveit, and A.B. Vardøy: <i>Metall. Mater. Trans. A</i>, 2015, vol. 46A, pp. 5266–74.</p> <p>69. S.A. Paknejad and S.H. Mannan: <i>Microelectron. Reliab.</i>, 2017, vol. 70, pp. 1–11.</p> <p>70. K.S. Siow: <i>J. Electron. Mater.</i>, 2014, vol. 43, pp. 947–61.</p>	<p>658 659 660 661 662 663 664 665 666 667 668 669 670 671 672 673</p>
--	--	---	--

**Publisher's Note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

UNCORRECTED PROOF

	Journal : <b>MMTA</b>	Dispatch : <b>1-7-2019</b>	Pages : <b>10</b>
	PIPS No. : <b>5356</b>	<input type="checkbox"/> LE	<input type="checkbox"/> TYPESET
	MS Code :	<input type="checkbox"/> CP	<input type="checkbox"/> DISK