VACUUM BRAZING

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- 1. Brazing amongst joining processes
- 2. Physical principles of (vacuum) brazing
- 3. Fabrication cycle of vacuum brazed assemblies
- 4. Equipment and technical requirements
- 5. CERN-made examples

6. Conclusion





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Metal Joining at CERN – The Forming and Welding-Section (EN-MME-FW)



LHC-Magnets Interconnection



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Current Leads for superconducting magnets



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RF-Couplers of SPS-Accelerating Cavities



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Metal Joining at CERN – The Forming and Welding-Section (EN-MME-FW)



LHC Accelerating RF-Cavities



The terms **Brazing** and Soldering contain a group of welding processes, where components are joined by a filler metal whose liquidus temperature is below the solidus/melting temperatures of the components' materials.





Brazing Processes Classification

(ISO 4063 - "Welding and allied processes - Nomenclature of processes and reference numbers")



- 1 Arc Welding
- 2 Resistance Welding
- 3 Gas Welding
- 4 Welding with pressure
- 5 Beam Welding
- 7 Other welding processes
- 8 Cutting and gouging

9 Brazing, soldering and braze welding

91	Brazing with local heating	
92	Brazing with global heating	_
93	Other brazing processes	
94	Soldering with local heating	
95	Soldering with global heating	
96	Other soldering processes	
97	Weld brazing	

921	Furnace brazing	
922	Vacuum brazing	
923	Dip-bath brazing	
924	Salt-bath brazing	
925	Flux-bath brazing	
926	Immersion brazing	









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Physical principles of (vacuum) brazing

Objective:

Bonding interfaces of base materials with a filler metal

Every brazing process has to go through following steps:

- 1. Liquification of the filler metal
- 2. Penetration of the base materials' clearance by the liquid filler metal
- 3. Creating adhesion between filler and base materials
- 4. Solification of the filler metal







Physical principles of (vacuum) brazing

1. Liquification of the brazing filler metal (BFM)



Pure metals:

Concrete melting point (e.g. Silver: 962°C)

Mixture/Alloys (binary, tertiary,...):

- Range of fusion T_{solidus}/T_{liquidus}
- Eutectic reaction T_{melt}
- (Peritectic systems, intermetallic reactions, ...)





13

Θ

10-30°

≈40°

120-135°

120-140°

Physical principles of (vacuum) brazing

2. Penetration of the base materials' clearance by the liquid filler metal

Wetting of a solid phase by a liquid defined by surface energies γ \rightarrow Wetting angle γ_{SL} γ_{SG} γ_{SG} $\gamma = \frac{\Delta E}{\Delta A}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ Phase in contact with liquid metal (Solid/Liquid) $\rho = 90^{\circ}$ \rightarrow no wetting $\rho = 90^{\circ}$ \rightarrow de-wetting $\rho = 90^{\circ}$ \rightarrow de-wetting $\rho = 90^{\circ}$ \rightarrow de-wetting $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$ $\rho = 90^{\circ}$ $\gamma_{SG} - \gamma_{SL} = \gamma_{LG} \cos \theta$

Source: N. Eustathipoulos et al., Wettability at High Temperatures

Ceramics, e.g. oxides (AI_2O_3/Ag)

Problem: Most metals create an oxide-layer on their surface exposed to normal atmosphere, especially at higher temperatures! \rightarrow <u>metallic surface must be provided</u>



Low wetting angle \rightarrow good flow on base material

Liquid on a solid surface:

Physical principles of (vacuum) brazing





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Physical principles of (vacuum) brazing

2. Penetration of the base materials' clearance by the liquid filler metal

Liquid between solid surfaces:

Reduction of surface energy results in force

 \rightarrow Capillary pressure P_C

$$P_{C} = \frac{2(\gamma_{SL} - \gamma_{LG})}{e} = \frac{2\gamma_{LG}\cos\theta}{e}$$
$$P_{C} = g\rho_{BFM}h$$

Equilibrium for non-reactive wetting.

Typical values for molten metalls (e.g. AgCu28 on solid copper):

γ_{LG}≈0.9 J/m²; ρ=10⁴ kg/m³; *θ*≈10°

For *e*=200 µm → *h*≈0.1 m

For $e=30 \ \mu m \rightarrow h\approx 0.6 \ m$

The effective θ may increase due to surface roughness or local impurities
→ small gaps assure proper wetting
→ a minimum gap is needed to allow flow (defined by filler metals viscosity)

е

g

 $\theta > 90^{\circ}$

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Physical principles of (vacuum) brazing

3. Creating adhesion between filler and base materials

Metallic continuity to assure maximal possible joint strength

a) Solubility between filler material and base material

The solubility of the elements present at the provides the possibility that filler and base material can interact and create a "smooth" transition.

Not or poorly soluble metals will result in distinct, separate phases with the phase boundary as joint layer. \rightarrow weak mechanical strength







Physical principles of (vacuum) brazing

- 3. Creating adhesion between filler and base materials
- 4. Solidification of the filler metal

Metallic continuity to assure maximal possible joint strength

b) Diffusion to create a continuous transition

Depending on the filler and base material combination the components exchange material through diffusion.

- \rightarrow Continuous material transition with no discrete weak interface
- \rightarrow Dwell for a certain period in the range of the brazing temperature enhances diffusion and can improve the mechanical strength

Excessive diffusion: Porosity through Kirkendall-effect possible!







Physical principles of (vacuum) brazing

The principles of a brazing process are clear.

How does it work in real life?

• Heating of the workpiece

The volume of the brazed interface must be homogenously heated at brazing temperature:

Local heating: Torch/flame, inductive heating, laser, electron beam

Global heating: Immersion in molten liquid (salt, filler metal, flux), furnace heating (convection or radiation)

• Wetting of the base material with liquid filler metal

The filler metal (usually) only wets a metallic surface. Most metals possess an oxidised surface! Heating under exposure of oxygen may furtherly thicken the oxide-layer.

- Using flux agent to remove oxides by chemical/physical dilution during brazing. Mostly chemical compounds with halogen-atoms (F, Cl)
- → Commonly used for manual and automatic flame brazing. Parts need to be cleaned thoroughly afterwards as flux residuals are corrosive.

Brazing under inert or reducing atmosphere can provide a clean, metallic surface for brazing. Interaction of applied gases with materials must be considered.

Or it is done under vacuum!



Physical principles of vacuum brazing

Heating up metals under vacuum

Residual pressures with high vacuum systems in the range of 10^{-6} mbar $\rightarrow 10^{-9}$ (1 ppb) of atmosphere No further oxidation during heating/brazing cycle!

Thermodynamic stability of metal oxides:



 $\frac{2x}{y}M + O_2 \xrightarrow{\Delta G} \frac{2}{y}M_{\chi}O_y$

$$\Delta G = \Delta H - T \Delta S$$

q.: $\Delta G - RT \ln(\frac{p_{O_2}}{p_0}) = 0$

Oxides of e.g. Cu, Fe, Ni can be reduced at brazing temperatures >700°C and HVconditions!*

*kinetics not taken into account



Physical principles of vacuum brazing

Main considerations of vacuum Brazing

Metals with an initially thin oxide scale (prepared by mechanical/chemical cleaning) can be wetted under vacuum through

- mechanical expulsion by wetting/capillary force
- thermodynamic reduction through high T and low p_{O2}

The stability of formed oxides on metallic surfaces defines in first line the brazeability under vacuum:

Brazeability	Nature of Oxide	Example
good/excellent	Low enthalpy, less stable at HT	Ag, Cu, Ni
fair/difficult	Stable, slowly forming/thin scale	Cr, Nb, Ti
Very difficult	Stable, fast forming/thick scale	Al, Mg, Be









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Steps to assemblies joined by vacuum brazing







Compatibility with vacuum brazing process

- Brazeability of metallic surface
- Compatibility with vacuum heat treatment to brazing temperatures Vapour pressure, annealing
- Mismatch of coefficient of thermal expansion for different materials Brazing gap, residual stress



Allowing correct assembly and proper brazing gaps

• Vacuum brazing executed in a furnace chamber:

Stable assembly configurations

- \rightarrow Positioning and loading of assemblies
- → Alignment features











Fabrication of correct pieces

• Sourcing of raw material – Identification, certification





Residual stresses – raw state, cold working during machining process

As the brazing cycle represents a heat treatment, stress relief can cause undesired deformations

 \rightarrow Intermediate heat treatments for vacuum brazed components often required



Stress relief of copper under air (200-250 °C)



Stress relief of spin-formed FeNiCo-parts under vacuum (850°C)

Important step: The right surface treatment

- Degreasing/cleaning of all surface contaminants (oil and other hydrocarbons)
- Pickling to remove oxide scale and keep oxidation as little as possible
- Plating of certain surfaces to improve/enable wetting during brazing

Ni-layer on (stainless) steels

Metallisation layers on non-metallic/ceramic materials allow joining by brazing



curtesy of M. Mayer (CERN)

Vith the



Adequate assembly for furnace

- Alignment and charging of brazed interfaces maintained through the whole brazing process
- Stable placement of filler metal (wires, foils, powder/paste)
- Support structures and tooling must be compatible with heat treatment and thermal expansion of part







The actual brazing happens

- Pump-down of vacuum furnace to HV (<10⁻⁵ mbar)
- Heating cycle varies with filler metal, thermal inertia and diffusivity of load, material requirements
- Usual cycle duration including cooling >12 h





Stainless steel-copper assembly with Ag86.5Cu26.5Pd5 (ISO 17672-Pd 287)

T_{solidus} ≈ 810°C

Load thermocouple inserted in one of the flanges



Control of brazed assemblies

according to requirements (e.g. qualification according to EN 13134/ISO 13585)

- Visual control
- Leak tests
- Metrology
- Destructive Testing: Mechanical Testing, Metallographic Cut
- Non Destructive Testing: Ultrasound Inspection; X-Ray/µCT for small assemblies (material/spacial resolution)





... and what can go wrong:

Most appearing defects of vacuum brazed joints

Cavities/pores •

Lack of penetration •

Excess of filler metal •

Erosion/alloying by filler metal •







Illustrations from EN ISO 18279:2003





Microgrpahs with the curtesy of M. Meyer, M. Crouvizier





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Vacuum Furnace Chamber

Heating source:

- Resistors
- Induction
- (Infrared)

 $\frac{dQ_{net}}{dt} \approx \epsilon_1 \sigma \left(T_2^{4} - T_1^{4} \right)$

Hot zone/resistors made from refractory metals mainly

Insulation by thermal screens

Water-cooled vessels

 T_{max} (Mo-hot zone): 1300-1600°C Power rating (hot zone ≈ 700 I): 200 kW Usual temperature uniformity ±5-10°C (>600°C)





Vacuum Group

Pumping Group

Primary/Foreline:

Mechanical Pumps (rotary vane/roots booster)

 \rightarrow 10⁻³ mbar

High Vacuum Pump:

 $\begin{array}{lll} \text{Oil-Diffusion} & (p_{\min} \approx 10^{\text{-7}} \text{ mbar}), \ \text{S}_{\text{pump}} \uparrow \\ \text{Turbomolecular} & (p_{\min} \approx 10^{\text{-9}} \text{ mbar}), \ \text{S}_{\text{pump}} \rightarrow \\ \text{Cryopump} & (p_{\min} \approx 10^{\text{-9}} \text{ mbar}), \ \text{S}_{\text{pump}} \uparrow \uparrow \end{array}$







Vacuum Group

Generation of Vacuum

Vacuum Leak tightness: $10^{-6}...10^{-3}$ mbar/(I×s) $p_{t\to\infty} = \frac{Q_{leak}}{S_{pump}}$

Cleanliness (hydrocarbons, volatile elements)

Compression rates for molecular pumps:

 $\begin{array}{ll} \mathsf{N}_2/\mathsf{O}_2/\mathsf{Ar} & \approx 10^8 \\ \mathsf{He} & \approx 5 \times 10^5 \\ \mathsf{H}_2 & \approx 5 \times 10^4 \end{array}$

 \rightarrow predominantely H2 present in residual gas





Control and Automation

Automation per PLC Pumping sequence, Heating cycle...



Recording of process parameters Pressure readings, furnace-/pump-/load-temperatures



Brazing surveillance Load thermocouples, furnace windows (>700°C)







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Joining of dissimilar materials

Stainless steel - Copper



Transitions from ss-flanges to copper for vacuum chamber, RF-cavities,...

Septum coils for kicker magnets. Stainless steelcapillaries brazed into conductor



Usual filler metals:

- Ag72/Cu28 (eutectic, T_{braze}: 780°C)
- Ag68/Cu27/Pd5 (T_{braze}: ca. 815°C)
- Ag58/Cu32/Pd10 (T_{braze}: ca. 855°C)...





Joining of dissimilar materials

Stainless steel – Niobium



Transitions from ss-flanges to Nb for Superconducting RF-cavities,...

Lower membrane of QPR-Cavity

Joining of two materials with a significant mismatch in CTE! (AISI 316 ≈18×10⁻⁶ K⁻¹; Nb ≈5×10⁻⁶ K⁻¹) Procedure established by CERN around 30 years ago.

Filler metal:

• Pure Cu (UNS C10100), T_{melt}: 1084°C

Joining of dissimilar materials

Stainless steel – Titanium



Transitions from ss-tubes to Ti (cp) for He-tank of SRF-cavities

Joining of two materials with mismatch in CTE! (AISI 316 ≈18×10⁻⁶ K⁻¹; Ti ≈9×10⁻⁶ K⁻¹)

Filler metal:

• Ag88Pd9Ga9, T_{liq}: 880°C – Contains no copper that would form very brittle Ti_xCu_v-intermetallics



Joining of dissimilar materials

Metal – Ceramic Transitions







RF-window for CRAB SRF-Cavity $Ti/Al_2O_3/Cu$

Insulator in chamber for BCT FeNiCo/Al $_2O_3$

Active Brazing of FeNiCo to AIN

Metals providing low CTE-mismatch with respect to corresponding ceramic (FeNiCo, Ti, Mo, Nb...)

Filler metal:

- Ag72/Cu28 and derivates, T_{braze}: 780-850°C (for metallised ceramics)
- Active Brazing with active brazing filler metals, e.g. Ag63Cu35.25Ti1.75

Joining of dissimilar materials

Metal – Ceramic Transitions



Non-ferromagnetic BCT-insulator $Al_2O_3/Cu/316L$

RF-feedthrough for HOM-couplers Cu/Al₂O₃/Cu/316L

Although high mismatch in CTE, Cu/Al_2O_3 -transitions feasible thanks to low yielding of copper

Filler metal:

• Metallised ceramic with additional Ag-plating creates in-situ a eutectic reaction (Liquid Transfer Brazing)









Assembly of RF-Cavities



RFQ for Linac4

Structure



Structure for e-acceleration for AWAKE

Precise assemblies – require often various machining and heat treatment steps Accuracy of assembly <20 µm over large dimensions





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Conclusions

When we use brazing – especially under vacuum?

- Joining of dissimilar materials
 - Different metalls with problematic metallurgic reactions during welding
 - Metal to ceramic-joints
- Joining of materials that are not possible or difficult to weld
 - · Copper and its alloys (high thermal conductivity, welding commonly performed, but delicate)
 - High melting refractory materials (high energy input, practically only welded by laser or electron beam)
 - Non-metallic materials (e.g. ceramics, carbon)
- Avoiding of welding shrinkage/fusion of base materials
 - Assemblies with tight tolerances
- Joints not accesible for welding
 - Assemblies with large areas to be joined
 - Complex joint geometries
 - Hidden joints



Conclusions

Some final words

- An evaluation of which joining/brazing process is the most adequate is important one vacuum brazing cycle can cost more than 1000 €
- Vacuum brazing is favorable for: Clean/UHV-components, large joint surfaces, precise assemblies of high quality, readily oxidizing/active base materials (Ti, Nb,...)
- The base materials can affect the joint design and the joint design can affect the base material (to be used)
- For a successful assembly, the brazing operator(s) should be involved from the design process on
- Each assembly needs specific process parameters/heating cycles
- Automated process: High reproducibility possible
- Taking all the steps into account, in will take days, or even weeks until an assembly is done by vacuum brazing – please be patient ☺☺☺





Thank you for your attention!







Thanks again for your attention!

